

DESCRIPTION

POLYHYDROXYALKANOATE HAVING VINYL GROUP, ESTER GROUP,
CARBOXYL GROUP AND SULFONIC ACID GROUP, AND

5 PRODUCTION METHOD THEREOF

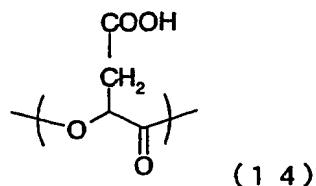
TECHNICAL FIELD

The present invention relates to a novel
polyhydroxyalkanoate and a production method thereof.

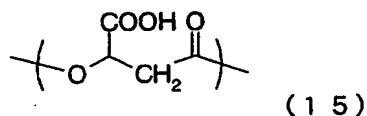
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BACKGROUND ART

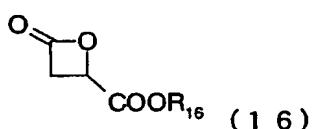
Biodegradable polymer materials are widely applied to medical materials, drug delivery systems, environmentally compatible materials, etc. In recent 15 years, new functions besides these are further demanded and various researches have been conducted. Particularly, introducing a chemically modifiable functional group into a molecule of polyhydroxyalkanoate represented by polylactic acid 20 is examined, and there are some reports about a compound to which a carboxyl group, a vinyl group or the like is introduced. For example, polymalic acid is known as a polyhydroxyalkanoate having a carboxyl group in the side chain. Among the polymers of this 25 polymalic acid, there have been known α -type compounds represented by the chemical formula (14) :



and β -type compounds represented by the chemical formula (15):

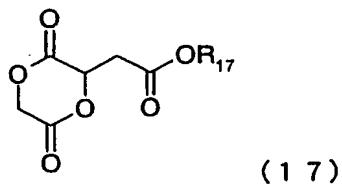


5 which are different in the way of polymer formation. Among these, as for the β -type polymalic acid and its copolymer, a polymer obtained by ring-opening polymerization of benzyl ester of β -malolactone represented by the chemical formula (16):



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(R₁₆: benzyl group) is disclosed in the specification of U.S. Patent No. 4265247. In addition, as for the α-type polymalic acid-glycolic acid copolymer and other copolymers containing hydroxyalkanoic acid including glycolic acid, polymers obtained by copolymerizing a 6-membered cyclic diester monomer represented by the chemical formula (17):



wherein R₁₇ represents a lower alkyl group such as a methyl group, an ethyl group, n-propyl group, an isopropyl group, and a t-butyl group and a benzyl group, etc., and a glycolide and lactide which are cyclic diesters, a lactone which is an ester by intramolecular ring closure reaction of ω -hydroxycarboxylic acid are disclosed in Japanese Patent Application Laid-Open No. H2-3415.

In addition, with regard to a polyhydroxyalkanoate having a carboxyl group on a side chain, it is disclosed in Macromolecules 2000, 33 (13), 4619-4627 that ring-opening polymerization of 7-oxo-4-oxepane carboxylate ester is conducted to prepare a polymer having an ester group on a side chain, which is further subjected to hydrocracking to prepare a polymer having carboxylic acid on a side chain. Biomacromolecules 2000, 1, 275 discloses a polymer to which benzyloxycarbonyl group is introduced at the α -position methylene group of a carbonyl group in the poly(ϵ -caprolactone) main chain by reacting poly(ϵ -caprolactone) with lithium diisopropylamide and further reacting with benzyl

chloroformate. *Macromolecular Bioscience* 2004, 4, 232 discloses a polymer to which (benzyloxycarbonyl)methyl group is introduced at the α -position methylene group of a carbonyl group in the 5 *polylactic acid* main chain by reacting *polylactic acid* with lithium diisopropyl amide and further reacting with benzyl bromoacetate.

With regard to a polyhydroxyalkanoate having a vinyl group on a side chain, *Polymeric Materials Science & Engineering* 2002, 87, 254 discloses a polymer obtained by ring-opening polymerization of α -allyl(δ -valerolactone). With regard to a polyhydroxyalkanoate having a vinyl group on a side chain, *Polymer Preprints* 2002, 43 (2), 727 also 10 discloses a polymer obtained by ring-opening polymerization of 3,6-diallyl-1,4-dioxane-2,5-dione which is a 6-membered ring diester monomer. 15

As mentioned above, there have been reported polymers having new functions imparted by further 20 introducing a functionality providing structure to a polyhydroxyalkanoate to which a chemically modifiable functional group has been introduced. In *International Journal of Biological Macromolecules* 25 (1999) 265, ring-opening polymerization of a cyclic 25 dimer of α -malic acid and glycolic acid gives a copolymer of α -type malic acid and glycolic acid, and a polyester having a carboxyl group on a side chain

is obtained by deprotecting the obtained polymer. It is described that carboxyl group of a side chain of this compound was subjected to chemical modification with a tripeptide, the obtained polymer was evaluated 5 about cellular adhesiveness and good results were obtained.

DISCLOSURE OF THE INVENTION

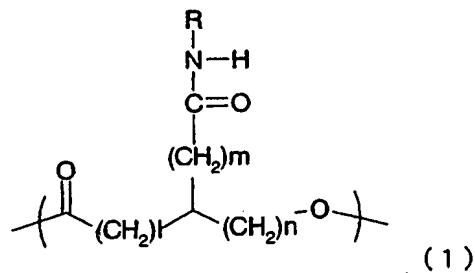
Although it is supposed to be possible to 10 impart a new functionality by introducing a unit having a carboxyl group or a unit having a vinyl group which are reactive functional groups into a molecule and carrying out chemical modification of the reactive functional group as mentioned above, 15 there have been few reports. Accordingly, the present invention provides a new polyhydroxyalkanoate having a reactive functional group in a molecule and a new polyhydroxyalkanoate having a new function imparted by performing chemical modification of the 20 polyhydroxyalkanoate having a reactive functional group and a production method thereof.

Therefore, the present inventors have conducted intensive studies aiming at developing a new polyhydroxyalkanoate having a reactive functional 25 group in a molecule and a new polyhydroxyalkanoate having a new function imparted by performing chemical modification of the polyhydroxyalkanoate having a

reactive functional group and consequently achieved the invention shown below.

Polyhydroxyalkanoates according to the present invention include the following.

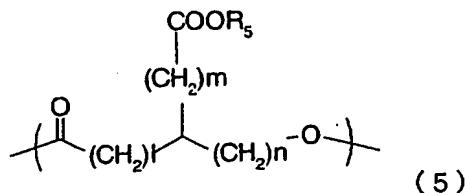
5 (1) A polyhydroxyalkanoate comprising one or more units represented by the chemical formula (1) in a molecule:



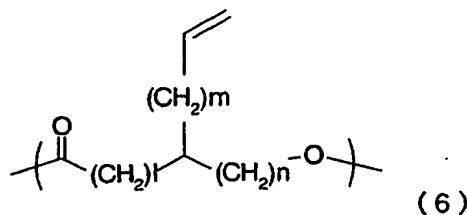
wherein R represents $-A_1-SO_2R_1$; R_1 is selected from 10 the group consisting of OH, a halogen atom, ONa, OK and OR_{1a}; R_{1a} and A₁ independently represent a group having a substituted or unsubstituted aliphatic hydrocarbon structure, a substituted or unsubstituted aromatic ring structure or a substituted or 15 unsubstituted heterocyclic structure, respectively; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and m is an integer selected from 0 to 8; and when two or more units are present, R, R₁, R_{1a}, A₁, and l, m and n mean as above 20 independently for every unit.

(2) A polyhydroxyalkanoate comprising one or more units represented by the chemical formula (5) in a

molecule:



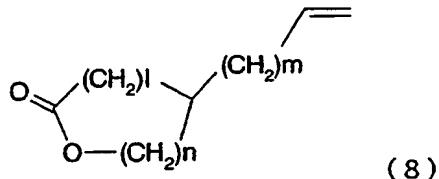
wherein R_5 is hydrogen, a salt forming group or R_{5a} ;
5 R_{5a} is a linear or branched alkyl group having 1 to 12
carbon atoms, an aralkyl group or a substituent
having a saccharide; l is an integer selected from 1
to 4, n is an integer selected from 1 to 4, m is an
integer selected from 0 to 8; and when l is 1, 3 and
4, n is an integer selected from 1 to 4, and m is an
integer selected from 0 to 8; and when l is 2 and n
10 is 1, 3 and 4, m is an integer selected from 0 to 8;
and when l is 2 and n is 2, m is an integer selected
from 1 to 8; and when l is 2, n is 2 and m is 0, R_{5a}
15 is a substituting group having a saccharide; and when
two or more units are present, R_5 , R_{5a} , and l , m and n
mean as above independently for every unit.



wherein l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and m is an integer selected from 0 to 8; and when two or more units are present, l, m, and n mean as above independently for every unit.

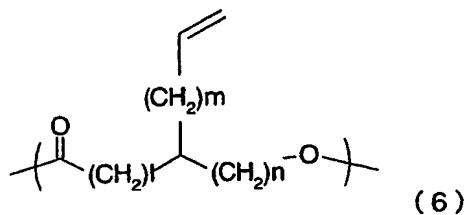
On the other hand, the production method of polyhydroxyalkanoate according to the present invention includes the following.

10 (A) A production method of polyhydroxyalkanoate represented by the chemical formula (6) having a step of polymerizing a compound having a unit represented by the chemical formula (8) in the presence of a catalyst.



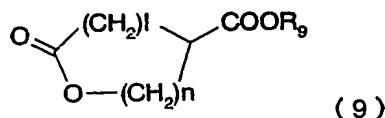
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wherein l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and m is an integer selected from 0 to 8.

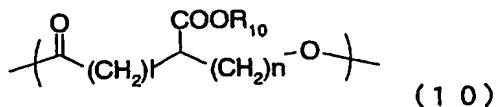


wherein l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and m is an integer selected from 0 to 8; and when two or more units are present, l, m and n mean as above independently for every unit.

(B) A production method of polyhydroxyalkanoate represented by the chemical formula (10) having a step of polymerizing a compound represented by the chemical formula (9) in the presence of a catalyst.

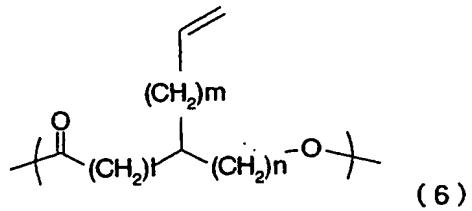


wherein R₉ is a substituent selected from a linear or branched alkyl group having 1 to 12 carbon atoms or an aralkyl group; l is an integer selected from 1 to 15 4, n is an integer selected from 1 to 4, and when l is 1, 3 or 4, n is an integer selected from 1 to 4, and when l is 2, n is 1, 3 or 4.)

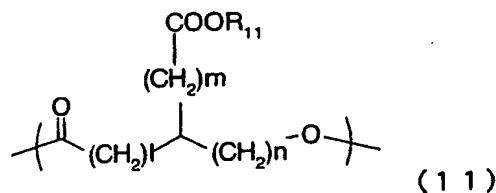


wherein R_{10} is a substituent selected from a linear or branched alkyl group having 1 to 12 carbon atoms or an aralkyl group; 1 is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and when 1 is 1, 3 or 4, n is an integer selected from 1 to 4, and when 1 is 2, n is 1, 3 or 4; and when two or more units are present, 1, n and R_{10} mean as above independently for every unit.

10 (C) A production method of polyhydroxyalkanoate
containing a unit represented by the chemical formula
(11) having a step of oxidizing a double bond portion
of polyhydroxyalkanoate containing a unit represented
by the chemical formula (6):

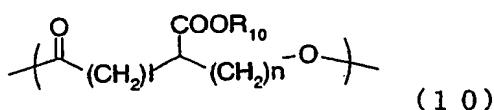


15 wherein l is an integer selected from 1 to 4, n is
an integer selected from 1 to 4, and m is an integer
selected from 0 to 8; and when two or more units are
present, l, m and n mean as above independently for
20 every unit.



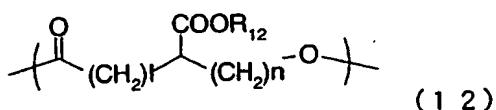
wherein R_{11} is hydrogen or a salt forming group; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and m is an integer selected from 0 to 8; and when two or more units are present, 5 l , m , n and R_{11} mean as above independently for every unit.

(D) A production method of polyhydroxyalkanoate containing a unit represented by the chemical formula 10 (12) having a step of hydrolyzing a polyhydroxyalkanoate containing a unit represented by the chemical formula (10) in the presence of acid or alkali, or subjecting a polyhydroxyalkanoate containing a unit represented by the chemical formula 15 (10) to hydrocracking including catalytic reduction:



wherein R_{10} is a substituent selected from a linear or branched alkyl group having 1 to 12 carbon atoms or an aralkyl group; l is an integer selected from 1 to 20 4, n is an integer selected from 1 to 4, and when 1

is 1, 3 or 4, n is an integer selected from 1 to 4, and when 1 is 2, n is 1, 3 or 4; and when two or more units are present, 1, n and R_{10} mean as above independently for every unit.

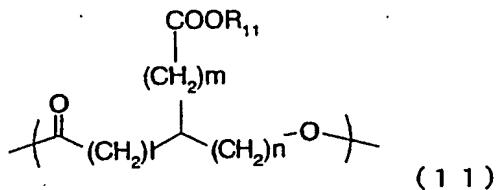


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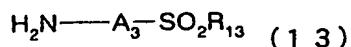
wherein R_{12} is hydrogen or a salt forming group; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and when l is 1, 3, and 4, n is an integer selected from 1 to 4, and when l is 2, n is 1, 3, and 4; and when two or more units are present, l , n and R_{12} mean as above independently for every unit.

(E) A production method of polyhydroxyalkanoate containing a unit represented by the chemical formula

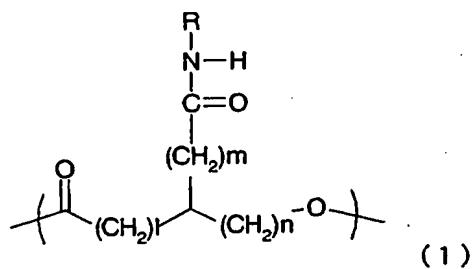
15 (1) having a step of subjecting a polyhydroxyalkanoate containing a unit represented by the chemical formula (11) and at least one amine compound represented by the chemical formula (13) to condensation reaction:



wherein R_{11} is hydrogen or a salt forming group; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and m is an integer selected from 0 to 8; and when two or more units are present, 5 l , m , n and R_{11} mean as above independently for every unit.



wherein R_{13} is selected from the group consisting of OH , a halogen atom, ONa , OK and OR_{13a} ; R_{13a} and A_3 are 10 independently selected from a group having a substituted or unsubstituted aliphatic hydrocarbon structure, a substituted or unsubstituted aromatic ring structure or a substituted or unsubstituted heterocyclic structure, respectively; and when two or 15 more units are present, R_{13} , R_{13a} and A_3 mean as above independently for every unit.



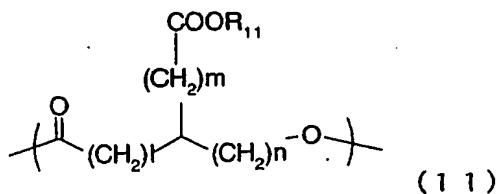
wherein R represents $-A_1-SO_2R_1$; R_1 is selected from the group consisting of OH , a halogen atom, ONa , OK

and OR_{1a} ; R_{1a} and A_1 independently represent a group having a substituted or unsubstituted aliphatic hydrocarbon structure, a substituted or unsubstituted aromatic ring structure or a substituted or 5 unsubstituted heterocyclic structure, respectively; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and m is an integer selected from 0 to 8; and when two or more units are present, R , R_1 , R_{1a} , A_1 , l , m and n mean as above independently 10 for every unit.

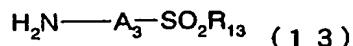
BEST MODE FOR CARRYING OUT THE INVENTION

Hereinbelow, the present invention is described in more detail referring to preferable embodiments.

15 The target polyhydroxyalkanoate represented by the chemical formula (1) in the present invention can be prepared by reacting polyhydroxyalkanoate containing a unit represented by the chemical formula (11) used as a starting material and at least one aminosulfonic 20 acid compound represented by the chemical formula (13).



wherein R_{11} is hydrogen or a salt forming group; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and m is an integer selected from 0 to 8; and when two or more units are present, 5 l , m , n and R_{11} mean as above independently for every unit.

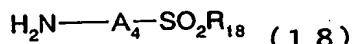


wherein R_{13} is selected from the group consisting of OH , a halogen atom, ONa , OK and OR_{13a} ; R_{13a} and A_3 are 10 independently selected from a group having a substituted or unsubstituted aliphatic hydrocarbon structure, a substituted or unsubstituted aromatic ring structure or a substituted or unsubstituted heterocyclic structure, respectively (here, R_{13a} is a 15 monovalent group having a structure selected from these and A_3 is a divalent group having a structure selected from these); and when two or more units are present, R_{13} , R_{13a} and A_3 mean as above independently for every unit.

20 More in detail, R_{13} is selected from the group consisting of OH , a halogen atom, ONa , OK and OR_{13a} . R_{13a} is a linear or branched alkyl group having 1 to 8 carbon atoms or a substituted or unsubstituted phenyl group. A_3 represents a group having a linear or 25 branched substituted or unsubstituted C_1 to C_8

alkylene group, a substituted or unsubstituted phenylene group, a substituted or unsubstituted naphthalene group or a substituted or unsubstituted group having a heterocyclic structure containing one 5 or more N, S and O. When A_3 is a cyclic structure, (a) unsubstituted ring(s) may be further condensed. Further, when two or more units are present, R_{13} , R_{13a} and A_3 mean as above independently for every unit.

In the case that A_3 is a linear and substituted 10 or unsubstituted alkylene group, aminosulfonic acid compounds represented by the following chemical formula (18) can be mentioned.

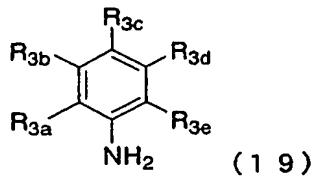


wherein R_{18} is selected from the group consisting of 15 OH, a halogen atom, ONa, OK and OR_{18a}; R_{18a} is a linear or branched alkyl group having 1 to 8 carbon atoms or a substituted or unsubstituted phenyl group; A_4 is a linear or branched substituted or unsubstituted C₁ to 20 C₈ alkylene group and when it is substituted, it may be substituted with an alkyl group having 1 to 20 carbon atoms, an alkoxy group having 1 to 20 carbon atoms, etc.

As a compound represented by the chemical formula (18), 2-aminoethanesulfonic acid (taurine), 25 3-aminopropanesulfonic acid, 4-aminobutanesulfonic

acid, 2-amino-2-methylpropanesulfonic acid, and alkaline metal salts and ester compounds thereof, etc. can be mentioned.

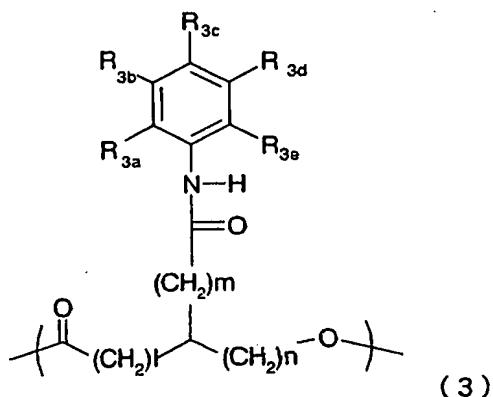
In the case that A_3 is a substituted or
5 unsubstituted phenylene group, aminosulfonic acid compounds represented by the following chemical formula (19) can be mentioned.



wherein R_{3a} , R_{3b} , R_{3c} , R_{3d} and R_{3e} are independently
10 selected from the group consisting of SO_2R_{3f} , wherein R_{3f} is selected from the group consisting of OH, a halogen atom, ONa, OK and OR_{3f1} wherein OR_{3f1} is a linear or branched alkyl group having 1 to 8 carbon atoms or a substituted or unsubstituted phenyl group,
15 a hydrogen atom, a halogen atom, an alkyl group having 1 to 20 carbon atoms, an alkoxy group having 1 to 20 carbon atoms, OH group and NH_2 group and NO_2 group, $COOR_{3g}$, wherein R_{3g} represents any of H atom, Na atom and K atom, an acetamide group, OPh group,
20 NPh group, CF_3 group, C_2F_5 or C_3F_7 group, respectively, wherein Ph represents a phenyl group, and at least one of the groups of these is SO_2R_{3f} ; and when two or more units are present, R_{3a} , R_{3b} , R_{3c} , R_{3d} , R_{3e} , R_{3f} , R_{3f1}

and R_{3g} mean as above independently for every unit.

By using the compound represented by the chemical formula (19), polyhydroxyalkanoate which contains in a molecule one or more units represented 5 by the chemical formula (3) can be obtained.

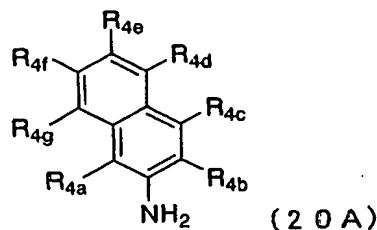


wherein R_{3a} , R_{3b} , R_{3c} , R_{3d} , R_{3e} , and l , n and m are similarly defined as the above.

As a compound represented by the chemical 10 formula (19), p-aminobenzenesulfonic acid (sulfanilic acid), m-aminobenzenesulfonic acid, o-aminobenzenesulfonic acid, m-toluidine-4-sulfonic acid, o-toluidine-4-sulfonic acid sodium salt, p-toluidine-2-sulfonic acid, 4-methoxyaniline-2-sulfonic acid, o-anisidine-5-sulfonic acid, p-anisidine-3-sulfonic acid, 3-nitroaniline-4-sulfonic acid, 2-nitroaniline-4-sulfonic acid sodium salt, 4-nitroaniline-2-sulfonic acid sodium salt, 1,5-dinitroaniline-4-sulfonic acid, 2-aminophenol-4-hydroxy-5-nitrobenzenesulfonic acid, 2,4-

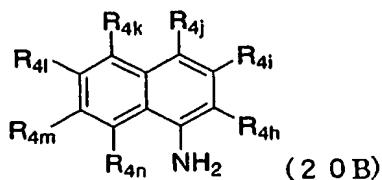
dimethylaniline-5-sulfonic acid sodium salt, 2,4-dimethylaniline-6-sulfonic acid, 3,4-dimethylaniline-5-sulfonic acid, 4-isopropylaniline-6-sulfonic acid, 4-trifluoromethylaniline-6-sulfonic acid, 3-carboxy-5 4-hydroxyaniline-5-sulfonic acid, 4-carboxyaniline-6-sulfonic acid and alkaline metal salts and ester compounds thereof, etc. can be mentioned.

In the case that A_3 is a substituted or unsubstituted naphthalene group, aminosulfonic acid 10 compounds represented by the following chemical formula (20A) or (20B) can be mentioned.



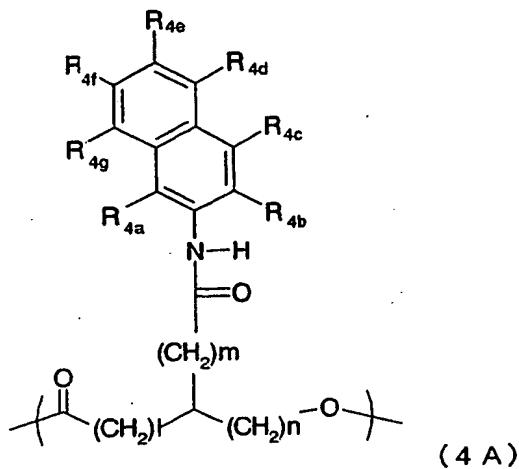
wherein R_{4a} , R_{4b} , R_{4c} , R_{4d} , R_{4e} , R_{4f} and R_{4g} are independently SO_2R_{4o} , wherein R_{4o} is selected from the 15 group consisting of OH, a halogen atom, ONa, OK and OR_{4o1} , wherein OR_{4o1} is a linear or branched alkyl group having 1 to 8 carbon atoms or a substituted or unsubstituted phenyl group; a hydrogen atom, a halogen atom, an alkyl group having 1 to 20 carbon atoms, an alkoxy group having 1 to 20 carbon atoms, 20 OH group, NH_2 group, NO_2 group, $COOR_{4p}$ group, wherein R_{4p} represents any of H atom, Na atom and K atom; an acetamide group, OPh group, $NHPh$ group, CF_3 group,

C_2F_5 group or C_3F_7 group, wherein Ph represents a phenyl group, respectively, and at least one of these groups is SO_2R_{40} .

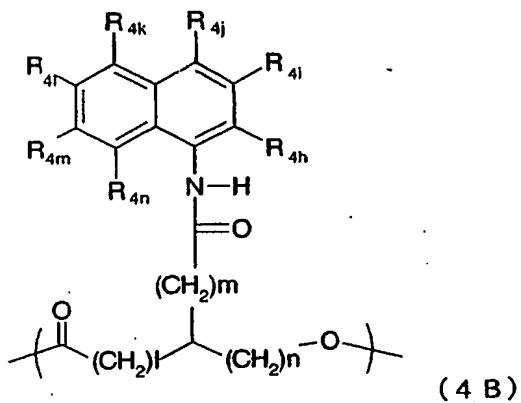


5 wherein R_{4h} , R_{4i} , R_{4j} , R_{4k} , R_{4l} , R_{4m} and R_{4n} are independently SO_2R_{40} wherein R_{40} is selected from the group consisting of OH, a halogen atom, ONa, OK and OR_{401} , wherein R_{401} is a linear or branched alkyl group having 1 to 8 carbon atoms or a substituted or 10 unsubstituted phenyl group; a hydrogen atom, a halogen atom, an alkyl group having 1 to 20 carbon atoms, an alkoxy group having 1 to 20 carbon atoms, OH group, NH_2 group, NO_2 group, $COOR_{4p}$ group, wherein R_{4p} represents any of H atom, Na atom and K atom; an 15 acetamide group, OPh group, NHPh group, CF_3 group, C_2F_5 group or C_3F_7 group, wherein Ph represents a phenyl group, respectively, and at least one of these groups is SO_2R_{40} .

Polyhydroxyalkanoate which contains one or more 20 units represented by the chemical formula (4A) or (4B) in a molecule can be obtained by using a compound represented by the chemical formula (20A) or (20B).



wherein R_{4a} , R_{4b} , R_{4c} , R_{4d} , R_{4e} , R_{4f} and R_{4g} and l , n and m are defined same as above.



5 wherein R_{4h} , R_{4i} , R_{4j} , R_{4k} , R_{4l} , R_{4m} and R_{4n} and l , n and m are defined same as above.

As a compound represented by the chemical formula (20A) or (20B), 1-naphthylamine 5-sulfonic acid, 1-naphthylamine 4-sulfonic acid, 1-naphthylamine 8-sulfonic acid, 2-naphthylamine 5-sulfonic acid, 1-naphthylamine 6-sulfonic acid, 1-

naphthylamine-7-sulfonic acid, 1-naphthylamine-2-ethoxy-6-sulfonic acid, 1-amino-2-naphthol-4-sulfonic acid, 6-amino-1-naphthol-3-sulfonic acid, 1-amino-8-naphthol-2,4-sulfonic acid monosodium salt, 1-amino-5-8-naphthol-3,6-disulfonic acid monosodium salt, and alkaline metal salts and ester compounds thereof, etc. can be mentioned.

In the case that A₃ is a substituted or unsubstituted group having a heterocyclic structure containing one or more N, S and O, the heterocyclic structure may be any one of pyridine ring, a piperazine ring, a furan ring, a thiol ring, etc. As a compound, sulfonic acid such as 2-aminopyridine-6-sulfonic acid and 2-aminopiperazine-6-sulfonic acid and alkaline metal salts and ester compounds thereof, etc. can be mentioned.

As a group which binds to a sulfonic acid through an ester bond in the case of sulfonic ester, a group having a substituted or unsubstituted aliphatic hydrocarbon structure, a substituted or unsubstituted aromatic ring structure or a substituted or unsubstituted heterocyclic structure can be mentioned as above. Furthermore, a linear or branched alkyl group having 1 to 8 carbon atoms, a substituted or unsubstituted phenyl group is preferable. From the viewpoints such as readiness of esterification, OCH₃, OC₂H₅, OC₆H₅, OC₃H₇, OC₄H₉,

$\text{OCH}(\text{CH}_3)_2$, $\text{OCH}_2\text{C}(\text{CH}_3)_3$, $\text{OC}(\text{CH}_3)_3$, etc. are more preferable.

(Production method of polyhydroxyalkanoate having a unit represented by the chemical formula (1))

5 A reaction of polyhydroxyalkanoate containing a unit represented by the chemical formula (11) with an aminosulfonic acid compound represented by the chemical formula (13) in the present invention is described in detail.

10 The amount of the compound represented by the chemical formula (13) used in the present invention is in the range of 0.1 to 50.0 times in mol, preferably 1.0 to 20.0 times in mol for a unit represented by the chemical formula (11) used as a 15 starting material.

There is a condensation reaction by thermal dehydration etc. as a method of generating an amide bond from the carboxylic acid and amine of the present invention. Particularly, a method of activating a carboxylic acid moiety by an activating agent to generate an active acyl intermediate and then reacting it with an amine is effective from the viewpoint of such mild reaction conditions so that ester bonds in the polymer main chain may not be cleaved. As an active acyl intermediate, an acid halide, an acid anhydride, active ester, etc. can be mentioned. Particularly, a method of using

condensing agent and forming an amide bond all over the same reaction space is preferable from the viewpoint of simplification of productive process. If necessary, it is also possible to perform a 5 condensation reaction with an amine after isolating as an acid halide.

As a usable condensing agent, a phosphate condensing agent used for polycondensation of aromatic polyamide, a carbodiimide condensing agent 10 used for peptide synthesis, an acid chloride condensing agent, etc. can be suitably selected based on the combination of the compounds of chemical formulas (13) and (11).

As the phosphate condensing agent, a 15 phosphorous acid ester condensing agent, a phosphorus chloride condensing agent, a phosphoric acid anhydride condensing agent, a phosphoric acid ester condensing agent, a phosphoric acid amide condensing agent, etc. can be mentioned.

20 It is possible to use a condensing agent such as a phosphorous acid ester in the reaction of the present invention. Examples of phosphorous acid ester used on this occasion include triphenyl phosphite, diphenyl phosphite, tri-o-tolyl phosphite, 25 di-o-tolyl phosphite, tri-m-tolyl phosphite, di-m-tolyl phosphite, tri-p-tolyl phosphite, di-p-tolyl phosphite, di-o-chlorophenyl phosphite, tri-p-

chlorophenyl phosphite, di-p-chlorophenyl phosphite, trimethyl phosphite, triethyl phosphite, etc.

Particularly, triphenyl phosphite is used preferably.

A metal salt such as lithium chloride and calcium

5. chloride may be also added for the improvement of properties such as solubility of and reactivity the polymer.

As the carbodiimide condensing agent,

dicyclohexylcarbodiimide (DCC), N-ethyl-N'-3-

10 dimethylaminopropylcarbodiimide (EDC=WSCI), diisopropylcarbodiimide (DIPC), etc. can be mentioned. N-hydroxysuccinimide (HONSu), 1-hydroxybenzotriazole (HOBt) or 3-hydroxy-4-oxo-3,4-dihydro-1,2,3-benzotriazine (HOObt), etc. may be used in

15 combination with DCC or WSCI.

The amount of the condensing agent used is in the range of 0.1 to 50 times in mol, preferably 1.0 to 20 times in mol for a unit represented by the chemical formula (11).

20 A solvent can be used, if needed, in the reaction of the present invention. The solvents to be used include hydrocarbons such as hexane, cyclohexane and heptane, ketones such as acetone and methyl ethyl ketone, ethers such as dimethyl ether, diethyl ether, tetrahydrofuran, halogenated hydrocarbons such as dichloromethane, chloroform, carbon tetrachloride, dichloroethane and

trichloroethane, aromatic hydrocarbons such as benzene and toluene, aprotic polar solvents such as N,N-dimethylformamide, dimethylsulfoxide, dimethylacetamide, and hexamethylphosphoramide, 5 pyridine, pyridine derivatives such as picoline, N-methylpyrrolidone, etc. Pyridine, N-methylpyrrolidone, etc. are used particularly preferably. The amount of the solvent used can be suitably determined depending on the start materials, 10 the kind of base, reaction conditions, etc. Although the reaction temperature is not particularly limited in the method of the present invention, it is usually a temperature in the range of -20°C to the boiling point of the solvent. However, it is preferable to 15 conduct reaction at the optimal temperature for the condensing agent to be used.

In the method of the present invention, the range of reaction time is usually 1 to 48 hours. Particularly, 1 to 10 hours is preferable.

20 In the present invention, collection and purification of the target polyhydroxyalkanoate from a reaction liquid containing the thus generated polyhydroxyalkanoate having a unit represented by the chemical formula (1) can be carried out by a normal method such as distillation. Alternatively, the 25 target polyhydroxyalkanoate represented by the chemical formula (1) can be precipitated by uniformly

mixing a solvent such as water, alcohols such as methanol and ethanol, ethers such as dimethyl ether, diethyl ether and tetrahydrofuran with the reaction liquid, and thereby collecting the same.

5 Polyhydroxyalkanoate having a unit represented by the chemical formula (1) obtained here can be isolated and purified if required. As this isolating and purifying method, there is particularly no restriction and the method of re-precipitating using
10 10 a solvent insoluble to polyhydroxyalkanoate having a unit represented by the chemical formula (1), method by column chromatography, dialyzing method, etc. can be used.

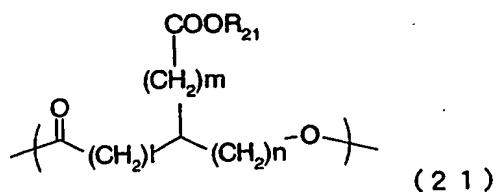
As another production method of the present
15 invention, when R moiety in the chemical formula (1) is $-A_1-SO_3H$, there is a method of performing methyl esterification to convert the R moiety in chemical formula (1) to $-A_1-SO_3CH_3$ using a methyl esterification agent after condensation reaction with
20 an amine. As a methyl esterification agent, those used for methyl esterification of a fatty acid in gas chromatography analysis can be used. As a methyl esterification method, an acid catalyst method such as hydrochloric acid-methanol method, boron trifluoride-methanol method and sulfuric acid-methanol method, a base catalyst method such as sodium methoxide method, tetramethylguanidine method

and trimethylsilyldiazomethane method, etc., can be mentioned. Particularly, trimethylsilyldiazomethane method is preferable since methylation is possible under mild conditions.

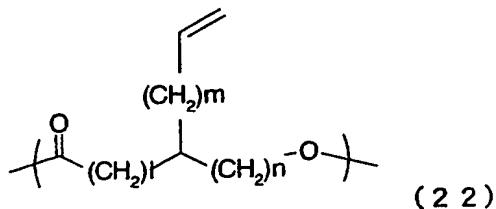
5 As for the solvent used in the reaction of the present invention, hydrocarbons such as hexane, cyclohexane and heptane, alcohols such as methanol and ethanol, halogenated hydrocarbons such as dichloromethane, chloroform, carbon tetrachloride, 10 dichloroethane, and trichloroethane, aromatic hydrocarbons such as benzene and toluene, etc. can be mentioned. Halogenated hydrocarbons are used particularly preferably. The amount of the solvent used can be suitably determined depending on the 15 starting materials, reaction conditions, etc. Although the reaction temperature is not particularly limited in the method of the present invention, it is usually a temperature in the range of -20°C to 30°C. However, it is preferable to conduct reaction at the 20 optimal temperature for the condensing agent and reagent to be used.

In the meantime, among the polyhydroxyalkanoate having a unit represented by the chemical formula (5) of the present invention, a polyhydroxyalkanoate 25 having a unit represented by the chemical formula (21) can be prepared by using a polyhydroxyalkanoate having a unit represented by the chemical formula

(22) as a starting material and oxidizing a double bond moiety in the side chain of the polyhydroxyalkanoate.



5 wherein R_{21} is hydrogen or a salt forming group; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, m is an integer selected from 0 to 8, when l is 1, 3, and 4, n is an integer selected from 1 to 4, and m is an integer selected from 0 to 10; and when l is 2 and n is 1, 3, and 4, m is an integer selected from 0 to 8; and when l is 2 and n is 2, m is an integer selected from 1 to 8; and when two or more units are present, R_{21} , l , m , and n mean as above independently for every unit.



wherein l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, m is an integer selected from 0 to 8, when l is 1, 3, and 4, n is an

integer selected from 1 to 4, and m is an integer selected from 0 to 8; and when l is 2 and n is 1, 3, and 4, m is an integer selected from 0 to 8; and when l is 2 and n is 2, m is an integer selected from 1 to 5 8; and when two or more units are present, l, m and n mean as above independently for every unit.

As a method of carrying out oxidative cleavage of a carbon-carbon double bond by an oxidizing agent as above to obtain a carboxylic acid, for example, a 10 method using a permanganic acid salt (J. Chem. Soc., Perkin. Trans. 1, 806 (1973)), a method using a dichromate (Org. Synth., 4, 698 (1963)), a method using a periodic acid salt (J. Org. Chem., 46, 19 (1981)), a method using nitric acid (Japanese Patent 15 Application Laid-Open No. S59-190945), a method using ozone (J. Am. Chem. Soc., 81, 4273 (1959)) etc. are known. Macromolecular chemistry, 4, 289-293 (2001) also reports a method of obtaining a carboxylic acid by reacting a carbon-carbon double bond at the end of 20 the side chain of polyhydroxyalkanoate produced by microorganisms using potassium permanganate as an oxidizing agent under acidic conditions. The same method can be used also in the present invention.

As a permanganic acid salt used as an oxidizing 25 agent, potassium permanganate is commonly used. The amount of a permanganic acid salt used may be usually 1 mol equivalent or more and preferably 2 to 10 mol

equivalents per one mol of the unit represented by the chemical formula (22), since the oxidative cleavage reaction is a stoichiometric reaction. In order to adjust the reaction system under acidic 5 conditions, various kinds of inorganic acids and organic acids such as sulfuric acid, hydrochloric acid, acetic acid and nitric acid are usually used. However, when acids such as sulfuric acid, nitric acid, hydrochloric acid and the like are used, ester 10 bond in the main chain may be cleaved, which may cause a decrease in the molecular weight. Therefore, it is preferable to use acetic acid. The amount of acid used is usually 0.2 to 2000 mol equivalent and preferably 0.4 to 1000 mol equivalent per one mol of 15 the unit represented by the chemical formula (22). When it is not less than 0.2 mol equivalent, preferable yield will be resulted, and when it is not more than 2000 mol equivalent, by-product material by decomposition by acid can be reduced, and therefore, 20 it is preferable to adjust it within the limits as defined above. Moreover, crown-ethers can be used in order to promote the reaction. In this case, crown-ether and permanganic acid salt form a complex, and result in an increased reaction activity. As a 25 crown-ether, dibenzo-18-crown-6-ether, dicyclo-18-crown-6-ether and 18-crown-6-ether are generally used. It is usually preferable to use the crown-ether in an

amount of usually 0.005 to 2.0 mol equivalent, and preferably 0.01 to 1.5 mol equivalent per one mol of permanganic acid salt.

The solvent used in the oxidation reaction of 5 the present invention is not particularly limited, as long as it is a solvent inactive in the reaction, and, for example, water, acetone; ethers such as tetrahydrofuran and dioxane; aromatic hydrocarbons such as benzene; aliphatic hydrocarbons such as 10 hexane and heptane; halogenated hydrocarbons such as methyl chloride, dichloromethane and chloroform can be used. Among these solvents, if the solubility of polyhydroxyalkanoate is taken into consideration, halogenated hydrocarbons such as methyl chloride, 15 dichloromethane and chloroform and acetone are preferable.

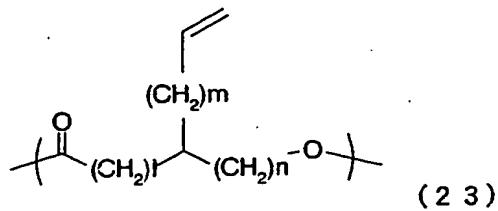
The polyhydroxyalkanoate having a unit represented by the chemical formula (22), permanganic acid salt and acid may be placed as a whole together 20 with a solvent from the beginning or each of them may be continuously or intermittently added to the system to proceed the reaction in oxidation reaction of the present invention. In addition, only permanganic acid salt may be previously dissolved or suspended in 25 a solvent, and subsequently polyhydroxyalkanoate and acid may be added to the system continuously or intermittently to proceed the reaction, or only

polyhydroxyalkanoate may be previously dissolved or suspended in a solvent, and subsequently permanganic acid salt and acid may be added to the system continuously or intermittently to proceed the reaction. Furthermore, polyhydroxyalkanoate and acid may be placed previously, and subsequently permanganic acid salt may be added to the system continuously or intermittently to proceed the reaction, or permanganic acid salt and acid are placed previously, and subsequently polyhydroxyalkanoate may be added to the system continuously or intermittently to proceed the reaction, or polyhydroxyalkanoate and permanganic acid salt are placed previously, and subsequently acid may be added to the system continuously or intermittently to proceed the reaction.

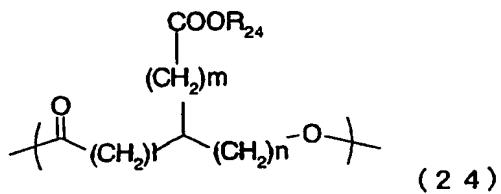
It is suitable to adjust the reaction temperature usually to -40 to 40°C, preferably -10 to 30°C. It is suitable to adjust the reaction time to usually 2 to 48 hours, although it is dependent on the stoichiometric ratio of the unit represented by the chemical formula (22) and permanganic acid salt and the reaction temperature.

In addition, polyhydroxyalkanoate containing a unit represented by the chemical formula (24) can be prepared by oxidation reaction from polyhydroxyalkanoate containing a unit represented by

the chemical formula (23) by the similar method as the above-described oxidation reaction. This production method is a novel production method.

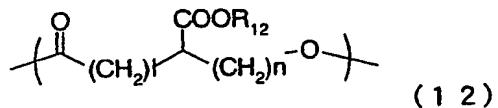


5 wherein l is 2, n is 2, and m is 0.

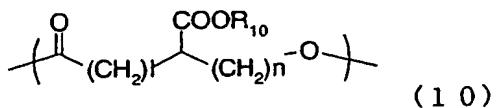


wherein R₂₄ is hydrogen or a salt forming group; l is 2, n is 2, and m is 0.

Among the polyhydroxyalkanoates having a unit 10 represented by the chemical formula (5), a polyhydroxyalkanoate having a unit represented by the chemical formula (12) can be also prepared by using a polyhydroxyalkanoate having a unit represented by the chemical formula (10) as a starting material by the 15 method of hydrolyzing the side chain ester moiety in the presence of an acid or alkali, or by the method of hydrocracking including catalytic reduction.



wherein R₁₂ is hydrogen or a salt forming group; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and when l is 1, 3, and 4, n is 5 an integer selected from 1 to 4, and when l is 2, n is 1, 3, and 4; and when two or more units are present, l, n and R₁₂ mean as above independently for every unit.



10 wherein R₁₀ is a substituent selected from a linear or branched alkyl group having 1 to 12 carbon atoms or an aralkyl group; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and when l is 1, 3 or 4, n is an integer selected from 1 to 4, 15 and when l is 2, n is 1, 3 or 4; and when two or more units are present, l, n and R₁₀ mean as above independently for every unit.

When a method of hydrolyzing in the presence of an acid or alkali is used, the method can be 20 performed using a solution of an inorganic acid such as hydrochloric acid, sulfuric acid, nitric acid or

phosphoric acid, or an organic acid such as trifluoroacetic acid, trichloroacetic acid, p-toluenesulfonic acid and methanesulfonic acid in water or an organic solvent having affinity to water

5 such as methanol, ethanol, tetrahydrofuran, dioxane, dimethylformamide and dimethylsulfoxide, or alternatively using an aqueous solution of an aqueous caustic alkali such as sodium hydroxide and potassium hydroxide or an alkali carbonate such as

10 sodium carbonate and potassium carbonate or an alcoholic solution of metal alkoxide such as sodium methoxide and sodium ethoxide. It is suitable to adjust the reaction temperature usually to 0 to 40°C and preferable to 0 to 30°C. It is suitable to adjust

15 the reaction time usually to 0.5 to 48 hours.

However, when hydrolyzation is conducted with an acid or alkali, ester bond in the main chain may be cleaved in any case and may cause decrease in molecular weight.

20 When a method of obtaining a carboxylic acid using a method of hydrocracking including catalytic reduction is used, the method can be performed as follows. That is, hydrogen is made to act in the presence of a reduction catalyst under normal or

25 increased pressure in a proper solvent at a temperature from -20°C to the boiling point of the solvent used, preferably 0 to 50°C, and catalytic

reduction is performed. Examples of the solvent to be used include water, methanol, ethanol, propanol, hexafluoroisopropanol, ethyl acetate, diethyl ether, tetrahydrofuran, dioxane, benzene, toluene,

5 dimethylformamide, pyridine, N-methylpyrrolidone, etc. A mixed solvent of these solvents can also be used. As a reduction catalyst, catalysts such as palladium, platinum, rhodium, etc. by itself or supported on a carrier, or Raney nickel, can be used. It is

10 suitable to adjust the reaction time usually to 0.5 to 72 hours. The thus formed reaction liquid containing a polyhydroxyalkanoate having a unit represented by the chemical formula (10) is collected as a crude polymer by filtering off the catalyst and

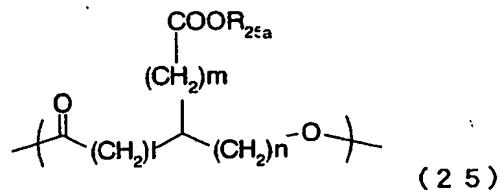
15 removing the solvent by distillation etc. The polyhydroxyalkanoate having a unit represented by the chemical formula (10) obtained here can be isolated and purified if required. The method of isolation and purification is not particularly limited and a

20 method of precipitating the polyhydroxyalkanoate having a unit represented by the chemical formula (10) using a solvent which does not dissolve the polyhydroxyalkanoate, a method using column chromatography, a dialyzing method, etc. can be used.

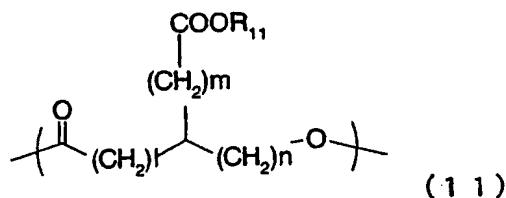
25 However, when catalytic reduction is used, ester bond in the main chain may be also cleaved and may cause a decrease in molecular weight.

In addition, among the polyhydroxyalkanoates having a unit represented by the chemical formula (5), a polyhydroxyalkanoate having a unit represented by the chemical formula (25) can be prepared by

5 esterifying a polyhydroxyalkanoate having a unit represented by the chemical formula (11) as a starting material using an esterification agent.



wherein R_{25a} is a linear or branched alkyl group having 1 to 12 carbon atoms or an aralkyl group or a substituent having a saccharide; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, m is an integer selected from 0 to 8, and when l is 1, 3, and 4, n is an integer selected from 1 to 4, and m is an integer selected from 0 to 8; and when l is 2 and n is 1, 3, and 4, m is an integer selected from 0 to 8; and when l is 2 and n is 2, m is an integer selected from 1 to 8; and when l is 2, n is 2 and m is 0, R_{25a} is a substituent having a saccharide; and when two or more units are present, R_{25a} , and l , m and n mean as above independently for every unit.

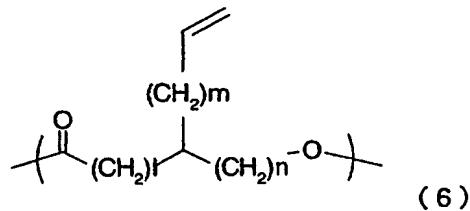


wherein R_{11} is hydrogen or a salt forming group; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and m is an integer selected from 0 to 8; and when two or more units are present, 5 l , m , n and R_{11} mean as above independently for every unit.

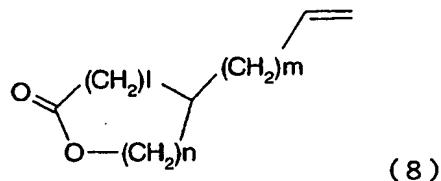
As an esterification agent used, diazomethane and DMF dimethyl acetal can be used. For example, 10 polyhydroxyalkanoate having a unit represented by the chemical formula (11) reacts easily with trimethylsilyldiazomethane, DMF dimethyl acetal, DMF diethyl acetal, DMF dipropyl acetal, DMF diisopropyl acetal, DMF-n-butyl acetal, DMF-tert-butyl acetal, or 15 DMF diisopentyl acetal, and gives corresponding esters. In addition, an esterified polyhydroxyalkanoate can be obtained by performing a reaction with an alcohol, for example, methanol, ethanol, propanol, isopropyl alcohol, butyl alcohol, 20 sec-butyl alcohol, tert-butyl alcohol, pentyl alcohol, neopentyl alcohol, hexyl alcohol, heptyl alcohol, octyl alcohol, nonyl alcohol, decyl alcohol and lauryl alcohol and saccharides for introducing a

group having a saccharide structure, for example, D-glucose, D-fructose and other saccharides following a method using an acid catalyst or a condensing agent such as DCC.

5 A polyhydroxyalkanoate represented by the chemical formula (6) of the present invention can be prepared by carrying out polymerization of an intramolecular closed ring compound of ω -hydroxycarboxylic acid represented by the chemical 10 formula (8) in the presence of a catalyst.



wherein l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and m is an integer selected from 0 to 8; and when two or more units are 15 present, l , m and n mean as above independently for every unit.



wherein l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and m is an integer

selected from 0 to 8.

In the production of a polyester containing a unit represented by the chemical formula (6) using an intramolecular closed ring compound of ω -5 hydroxycarboxylic acid represented by the chemical formula (8) of the present invention, polymerization method is not particularly limited and, for example, solution polymerization method, slurry polymerization method, mass polymerization method, etc. can be adopted. In addition, when a polymerization solvent is used, the solvent is not particularly limited and, for example, inert solvents such as aliphatic hydrocarbons and cyclic hydrocarbons having 5 to 18 carbon atoms, aromatic hydrocarbons having 6 to 20 carbon atoms, tetrahydrofuran, chloroform, α -dichlorobenzene, dioxane, etc. can be used. Conventional ring-opening polymerization catalysts can be used as a catalyst used for this polymerization. Examples thereof include tin dichloride, tin tetrachloride, stannous fluoride, stannous acetate, stannous stearate, stannous octanoate, stannous oxide, stannic oxide, and other tin salts. In addition, triethoxyaluminum, tri-n-propoxyaluminum, tri-iso-propoxyaluminum, tri-n-butoxyaluminum, tri-iso-butoxyaluminum, aluminium chloride, di-iso-propyl zinc, dimethylzinc, diethylzinc, zinc chloride, tetra-n-propoxytitanium,

tetra-n-butoxytitan, tetra-n-butoxytitan, tetra-t-butoxytitan, antimony trifluoride, lead oxide, lead stearate, titanium tetrachloride, boron trifluoride, boron trifluoride-ether complex, triethylamine, 5 tributylamine, etc. can be mentioned.

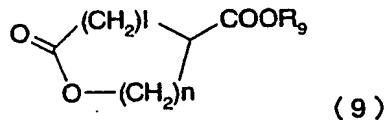
The amount of these catalysts to be used is in the range of 0.0001 to 10% by weight, and preferably in the range of 0.001 to 5% by weight to the total amount of the monomer compound.

10 In the present invention, conventional polymerization initiators can be used as a polymerization initiator on the occasion of ring-opening polymerization. Specifically, aliphatic alcohols may be any of mono-, di-, or polyhydric 15 alcohols and may be saturated or unsaturated alcohols. Specifically, monoalcohols such as methanol, ethanol, propanol, butanol, pentanol, hexanol, heptanol, nonanol, decanol, lauryl alcohol, myristic alcohol, cetyl alcohol, stearyl alcohol and p-tert-butylbenzyl 20 alcohol, dialcohols such as ethylene glycol, butanediol, hexanediol, nonanediol and tetramethylene glycol, polyhydric alcohols such as glycerol, sorbitol, xylitol, ribitol and erythritol, and methyl lactate, ethyl lactate, etc. can be used. Although 25 there are some differences according to the conditions depending on the alcohol used, these aliphatic alcohols can be usually used at a ratio of

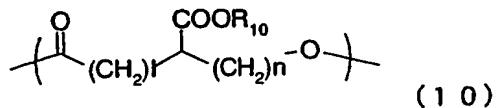
0.01 to 10% by weight to the total amount of the monomer. In the present invention, the temperature of ring-opening polymerization reaction is in the range of 25 to 200°C, preferably in the range of 50 to 200°C, more preferably in the range of 100 to 180°C.

In the present invention, the ring-opening polymerization reaction may be performed under inert atmosphere such as nitrogen and argon, either reduced or increased pressure, and a catalyst and an alcohol may be added sequentially.

In addition, the polyhydroxyalkanoate having a unit represented by the chemical formula (10) of the present invention can be prepared by carrying out polymerization of an intramolecular closed ring compound of ω -hydroxycarboxylic acid represented by the chemical formula (9) in the presence of a catalyst.



wherein R_9 is a substituent selected from a linear or branched alkyl group having 1 to 12 carbon atoms or an aralkyl group; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and when l is 1, 3 or 4, n is an integer selected from 1 to 4, and when l is 2, n is 1, 3 or 4.



wherein R_{10} is a substituent selected from a linear or branched alkyl group having 1 to 12 carbon atoms or an aralkyl group; l is an integer selected from 1 to 4, n is an integer selected from 1 to 4, and when l is 1, 3 or 4, n is an integer selected from 1 to 4, and when l is 2, n is 1, 3 or 4; and when two or more units are present, l and n mean as above independently for every unit.

10 In the production of a polyester containing a unit represented by the chemical formula (10) using an intramolecular closed ring compound of ω -hydroxycarboxylic acid represented by the chemical formula (9) of the present invention, polymerization 15 method is not particularly limited and, for example, solution polymerization method, slurry polymerization method, mass polymerization method, etc. can be adopted. In addition, when a polymerization solvent is used, the solvent is not particularly limited and, 20 for example, inert solvents such as aliphatic hydrocarbons and cyclic hydrocarbons having 5 to 18 carbon atoms, aromatic hydrocarbons having 6 to 20 carbon atoms, tetrahydrofuran, chloroform, α -dichlorobenzene, dioxane, etc. can be used.

In the present invention, conventional ring-opening polymerization catalysts can be used as a catalyst for polymerization. Examples thereof include tin dichloride, tin tetrachloride, stannous 5 fluoride, stannous acetate, stannous stearate, stannous octanoate, stannous oxide, stannic oxide, and other tin salts. In addition, triethoxyaluminum, tri-n-propoxy-aluminum, tri-iso-propoxyaluminum, tri-n-butoxyaluminum, tri-iso-butoxyaluminum, aluminium 10 chloride, di-iso-propyl zinc, dimethylzinc, diethylzinc, zinc chloride, tetra-n-propoxytitanium, tetra-n-butoxytitan, tetra-n-butoxytitan, tetra-t-butoxytitan, antimony trifluoride, lead oxide, lead stearate, titanium tetrachloride, boron trifluoride, 15 boron trifluoride-ether complex, triethylamine, tributylamine, etc. can be mentioned.

The amount of these catalysts to be used is in the range of 0.0001 to 10% by weight, and preferably in the range of 0.001 to 5% by weight to the total 20 amount of the monomer compound.

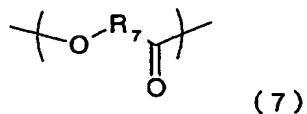
In the present invention, conventional polymerization initiators can be used as a polymerization initiator on the occasion of ring-opening polymerization. Specifically, aliphatic 25 alcohols may be any of mono-, di-, or polyhydric alcohols and may be saturated or unsaturated alcohols. Specifically, monoalcohols such as methanol, ethanol,

propanol, butanol, pentanol, hexanol, heptanol, nonanol, decanol, lauryl alcohol, myristic alcohol, cetyl alcohol, stearyl alcohol and p-tert-butylbenzyl alcohol, dialcohols such as ethylene glycol, 5 butanediol, hexanediol, nonanediol and tetramethylene glycol, polyhydric alcohols such as glycerol, sorbitol, xylitol, ribitol and erythritol, and methyl lactate, ethyl lactate, etc. can be used. Although there are some differences according to the 10 conditions depending on the alcohol used, these aliphatic alcohols can be usually used at a ratio of 0.01 to 10% by weight to the total amount of the monomer. In the present invention, the temperature of ring-opening polymerization reaction is in the 15 range of 25 to 200°C, preferably in the range of 50 to 200°C, more preferably in the range of 100 to 180°C.

In the present invention, the ring-opening polymerization reaction may be performed under inert atmosphere such as nitrogen and argon, either reduced 20 or increased pressure, and a catalyst and an alcohol may be added sequentially.

Although the polyhydroxyalkanoate of the present invention is mainly composed of the unit represented by the chemical formula (1), (5) or (6) 25 shown above, it may be also a copolymer to which a second ingredient etc. is copolymerized in order to change various physical properties such as mechanical

properties and decomposition characteristics. For example, a unit represented by the chemical formula (7) can be further contained in a molecule.



5 wherein R₇ is a linear or branched alkylene group having 1 to 11 carbon atoms, an alkyleneoxyalkylene group, wherein each alkylene group is independently an alkylene group having 1 to 2 carbon atoms, respectively, or an alkylidene group having 1 to 5 carbon atoms which may be substituted with aryl; and when two or more units are present, R₇ means as above 10 independently for every unit.

As an example as the second ingredient, a 15 cyclic diester of α -hydroxycarboxylic acid or a lactone which is an intramolecular closed ring compound of ω -hydroxycarboxylic acid can be copolymerized. Furthermore, specific examples of a cyclic diester of α -hydroxycarboxylic acid include intermolecular cyclic diesters such as glycolic acid, 20 lactic acid, α -hydroxybutyric acid, α -hydroxy-isobutyric acid, α -hydroxyvaleric acid, α -hydroxy-isovaleric acid, α -hydroxy- α -methylbutyric acid, α -hydroxycaproic acid, α -hydroxy-isocaproic acid, α -hydroxy- β -methylvaleric acid, α -hydroxyheptanoic acid,

mandelic acid, and β -phenyllactic acid. As for the compounds having an asymmetric carbon, any of L-compound, D-compound, racemate and meso compound can be used. The cyclic diester may be also formed from 5 different α -oxy acid molecules. Specifically, a cyclic diester between glycolic acid and lactic acid, and 3-methyl-2,5-diketo-1,4-dioxane etc. can be mentioned. Non-limiting examples of a lactone which is an intramolecular closed ring compound of ω -10 hydroxycarboxylic acid include β -propiolactone, β -butyrolactone, β -isovalerolactone, β -caprolactone, β -isocaprolactone, β -methyl- β -valerolactone, γ -butyrolactone, γ -valerolactone, δ -valerolactone, ϵ -caprolactone, 11-oxydecanoic acid lactone, p-15 dioxanone, 1,5-dioxepan-2-one.

Although the number average molecular weight of polyhydroxyalkanoate obtained by polymerization may vary depending on the conditions such as a type and amount of polymerization catalyst, polymerization 20 temperature and polymerization time, 1,000 to 1,000,000 is preferable.

The molecular weight of polyhydroxyalkanoate can be measured as a relative molecular weight as well as an absolute molecular weight. It can be 25 simply measured by GPC (gel permeation chromatography), etc. As a specific measuring method by GPC, the above-mentioned polyhydroxyalkanoate is

beforehand dissolved in a solvent which can dissolve the polyhydroxyalkanoate, and measurement is performed with a similar mobile phase. As a detector, a differential refraction detector (RI) and an 5 ultraviolet detector (UV) can be used depending on the polyhydroxyalkanoate to be measured. The molecular weight can be determined as a relatively compared value with a standard sample (polystyrene, polymethyl methacrylate, etc.). As a solvent, any 10 solvent which dissolves the polymer such as dimethylformamide (DMF), dimethylsulfoxide (DMSO), chloroform, tetrahydrofuran (THF), toluene, and hexafluoroisopropanol (HFIP) may be selected. In the case of a polar solvent, measurement can be also 15 performed while a salt is added.

In the present invention, it is preferable to use a polyhydroxyalkanoate as mentioned above which has a ratio (Mw/Mn) of the weight average molecular weight (Mw) to the number average molecular weight 20 (Mn) measured as mentioned above in the range of 1 to 10.

It should be noted that the reaction solvent, reaction temperature, reaction time, purifying method, etc. in the chemical reaction of the present 25 invention are not limited to the above-mentioned method.

Examples

Although the present invention is described in more detail referring to Examples given below and, the method of the present invention is not limited 5 only to these Examples.

First, materials used in the Examples are shown as Preparation Examples.

(Preparation Examples)

The materials used in the Examples can be 10 prepared by the following method.

(Preparation Example 1)

Production method of tetrahydro-4-(2-propenyl)-2H-pyran-2-one represented by the chemical formula (26) described in Examples 1 to 4.

15 Tetrahydro-4-(2-propenyl)-2H-pyran-2-one represented by the chemical formula (26) described in Examples 1 to 4 can be prepared by using the method described in Journal of American Chemical Society 2002, 124, 13171-13718 (compound (15) in this 20 document).

122 ml (244.0 mmol) of 2M allyl magnesium chloride-THF solution was placed in an eggplant flask and 500 ml of THF dissolving 27.5 g (122.1 mmol) of zinc bromide was added to this and agitated at 0°C. 25 This was placed in a nitrogen atmosphere and agitated at -78°C. Next, 53.0 g (487.8 mmol) of chlorotrimethylsilane was added, and 200 ml of THF

solution dissolving 10.0 g (101.9 mmol) of 3,4-dihydro-2H-pyran-2-one was further added slowly, and agitated at -78°C for 1 hour. 1N-hydrochloric acid was added to the reaction liquid after the reaction 5 ended and, then diethyl ether was added and the organic layer was extracted. The organic layer was collected, after it was washed with 1N-hydrochloric acid 3 times. The collected organic layer was dried over anhydrous sodium sulfate. After sodium sulfate 10 was removed, crude tetrahydro-4-(2-propenyl)-2H-pyran-2-one was collected by evaporating the solvent. Next, it was purified by silica gel column chromatography and 7.10 g of the target tetrahydro-4-(2-propenyl)-2H-pyran-2-one was obtained by 15 performing drying under reduced pressure. In order to identify the structure of the obtained compound, NMR analysis was performed on the following conditions.

<Measurement Apparatus> FT-NMR: Bruker DPX400
20 Resonant Frequency: ^1H = 400 MHz
<Measurement Conditions> Nuclide Measured: ^1H
Solvent Used: CDCl_3 ,
Measurement Temperature: room temperature
Consequently, it was confirmed that the 25 obtained compound was tetrahydro-4-(2-propenyl)-2H-pyran-2-one which was the object compound.
(Preparation Example 2)

Production method of tetrahydro-4-(3-butenyl)-2H-pyran-2-one represented by the chemical formula (31) described in Example 5.

7.40 g of the target tetrahydro-4-(3-butenyl)-2H-pyran-2-one was obtained by the same method as in Example 1 except that 122 ml (244.0 mmol) of 2M 3-butenyl magnesium bromide-THF solution was used in place of allyl magnesium chloride described in Example 1.

10 (Preparation Example 3)

Production method of 5-(2-propenyl)-2-oxepanone represented by the chemical formula (33) described in Example 6.

5-(2-propenyl)-2-oxepanone represented by chemical formula (33) described in Example 6 is a known compound and, for example, can be obtained by using the method described in Journal of American Chemical Society 1998, 120, 3541-3548 (compound (6e) in the above-mentioned document).

20 (Preparation Example 4)

Production method of 7-oxo-4-oxepane carboxylic acid phenylmethyl ester represented by the chemical formula (38) described in Example 8.

7-oxo-4-oxepane carboxylic acid phenylmethyl ester represented by the chemical formula (38) described in Example 8 can be obtained by using the method described in Macromolecules, 2000, 33, 4622

(compound (12) in this document).

The object compound can be obtained by four steps of reactions in more detail.

(Reaction 1)

5 To 280 ml of methylene chloride suspended solution containing 40.0 g (185.6 mmol) of pyridinium chlorochromate, 25 ml of methylene chloride solution containing 30.0 g (174.2 mmol) of ethyl 4-hydroxycyclohexane carbonate was added and agitated
10 at room temperature for 2 hours, and after that agitated at 45°C for 5 hours. The reaction solution was subjected to decantation after the reaction ended, and the residue was rinsed with diethyl ether. The reaction solution and the diethyl ether used for
15 rinsing were combined to one solution and purified by silica gel column chromatography followed by distillation under reduced pressure and thereby 29.5 g of ethyl 4-ketocyclohexane carbonate was obtained.

(Reaction 2)

20 2% sulfuric acid was added to 20.0 g (117.5 mmol) of ethyl 4-ketocyclohexane carbonate and agitated at 115°C for 4 hours. The reaction solution was extracted with diethyl ether after the reaction ended and dried over magnesium sulfate. Crude
25 product was obtained after the solvent was evaporated. By carrying out decompression distillation of the crude product, 13.5 g of ethyl 4-ketocyclohexane

carbonate was obtained.

(Reaction 3)

To an acetone solution containing 12.0 g (84.4 mmol) of 4-ketocyclohexane carboxylic acid and 4.7 g (33.8 mmol) of potassium carbonate, 17.3 g (101.3 mmol) of benzyl bromide was added at room temperature. 5 Next, this was agitated at 60°C for 3 hours. After the reaction ended, crude product was obtained, and after potassium carbonate was removed by filtration 10 and acetone was evaporated, the crude product was purified by silica gel column chromatography, and 16.5 g of 4-ketocyclohexane carboxylic acid phenylmethyl ester was obtained.

(Reaction 4)

15 To a chloroform solution containing 16.5 g (71.0 mmol) of benzyl 4-ketocyclohexane carbonate, a chloroform solution containing m-chloroperbenzoic acid was added at room temperature. This was agitated at 65°C for 2 hours. After the reaction 20 ended, the solution was filtered using Celite, and then an aqueous solution of sodium bicarbonate was added and the liquids were allowed to separate. After the organic layer was collected, it was dried over sodium sulfate. After the solvent was 25 evaporated, 14.7 g of the object compound, 7-oxo-4-oxepane carboxylic acid phenylmethyl ester represented by the chemical formula (38), was

obtained by purified by silica gel column chromatography followed by distillation under reduced pressure. In order to identify the structure of the obtained compound, NMR analysis was performed on the 5 same conditions as in Preparation Example 1, and consequently it was confirmed that the obtained compound was 7-oxo-4-oxepane carboxylic acid phenylmethyl ester represented by the chemical formula (38) which was the object compound.

10 (Preparation Example 5)

Production method of 7-oxo-3-oxepane carboxylic acid phenylmethyl ester represented by the chemical formula (35) described in Example 7.

7-oxo-3-oxepane carboxylic acid phenylmethyl ester represented by the chemical formula (35) described in Example 7 can be prepared by using the method described in Preparation Example 4. Specifically, the object compound can be obtained by performing (Reaction 3) and (Reaction 4) in the same 20 way except that 3-ketocyclohexanecarboxylic acid is used in place of 4-ketocyclohexane carboxylic acid in (Reaction 3) in Preparation Example 4.

6.0 g of the object compound, 7-oxo-3-oxepane carboxylic acid phenylmethyl ester represented by the 25 chemical formula (35), was obtained by performing a similar method in Preparation Example 4 except that 12.0 g (84.4 mmol) of 3-ketocyclohexanecarboxylic

acid was used in place of 4-ketocyclohexane carboxylic acid in Preparation Example 4.

(Preparation Example 6)

Production method of tetrahydro-6-oxo-2H-pyran-3-carboxylic acid phenylmethyl ester represented by the chemical formula (41) described in Example 9:

Tetrahydro-6-oxo-2H-pyran-3-carboxylic acid phenylmethyl ester represented by the chemical formula (41) described in Example 9 can be prepared by using the method described in Preparation Example 4. Specifically, the object compound can be obtained by performing (Reaction 3) and (Reaction 4) in the same way except that 3-ketocyclopentanecarboxylic acid is used in place of 4-ketocyclohexane carboxylic acid in (Reaction 3) in Preparation Example 4.

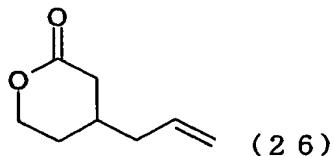
4.5 g of the object compound, tetrahydro-6-oxo-2H-pyran-3-carboxylic acid phenylmethyl ester represented by the chemical formula (41), was obtained by performing a similar method in Preparation Example 4 except that 10.8 g (84.4 mmol) of 3-ketocyclopentanecarboxylic acid was used in place of 4-ketocyclohexane carboxylic acid in Preparation Example 4.

Specific examples are shown below.

Example 1

(Synthesis of polyester using tetrahydro-4-(2-propenyl)-2H-pyran-2-one represented by chemical

formula (26))



1.40 g (0.01 mmol) of tetrahydro-4-(2-propenyl)-2H-pyran-2-one represented by the chemical formula (26), 4.0 ml of 0.01 M toluene solution of tin octylate (tin 2-ethylhexanoate), 4.0 ml of 0.01 M toluene solution of p-tert-butylbenzyl alcohol were placed in a polymerization ampoule and after performing drying under reduced pressure for 1 hour and nitrogen purge, it was heat-sealed under reduced pressure, and heated at 150°C to perform ring-opening polymerization. The reaction was ended after 12 hours and the reaction product was cooled. The obtained polymer was dissolved in chloroform and re-precipitated in methanol in an amount of 10 times of the chloroform required for dissolving the polymer. The precipitation was collected and 0.53 g of polymer was obtained by carrying out drying under reduced pressure. In order to identify the structure of the obtained polymer, NMR analysis was performed on the following conditions.

<Measurement Apparatus> FT-NMR: Bruker DPX400

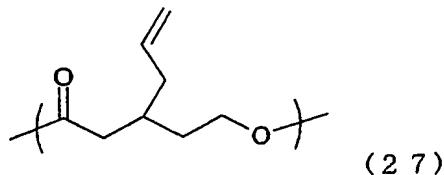
Resonant Frequency: ^1H = 400 MHz

<Measurement Conditions> Nuclide Measured: ^1H

Solvent Used: TMS/CDCl₃

Measurement Temperature: room temperature

Consequently, it was confirmed that the obtained compound was a polyhydroxyalkanoate which 5 consists of a unit represented by the following chemical formula (27).



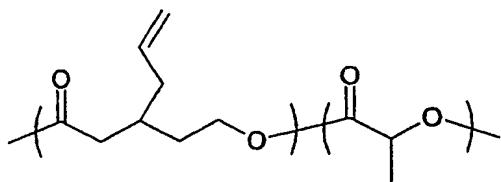
The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation 10 chromatography (GPC; TOSOH HLC-8220, column; TOSOH TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, Mn was 6,200, and weight average molecular weight, Mw was 8,100.

15 Example 2

(Synthesis of polyester using tetrahydro-4-(2-propenyl)-2H-pyran-2-one and L-lactide)

0.28g (2.0 mmol) of tetrahydro-4-(2-propenyl)-2H-pyran-2-one represented by the chemical formula 20 (26), 1.44 g (10.0 mmol) of L-lactide, 4.8 ml of 0.01 M toluene solution of tin octylate (tin 2-ethylhexanoate), 4.8 ml of 0.01 M toluene solution of p-tert-butylbenzyl alcohol were placed in a polymerization ampoule and after performing drying

under reduced pressure for 1 hour and nitrogen purge, it was heat-sealed under reduced pressure, and heated at 150°C to perform ring-opening polymerization. The reaction was ended after 12 hours and the reaction 5 product was cooled. The obtained polymer was dissolved in chloroform and re-precipitated in methanol in an amount of 10 times of the chloroform required for dissolving the polymer. The precipitation was collected and 1.19 g of polymer was 10 obtained by carrying out drying under reduced pressure. In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a 15 polyhydroxyalkanoate copolymer which contains a unit represented by the following chemical formula (28) as a monomer unit. It was also confirmed that the ratios of monomer units were 7 mol% for A unit and 93 mol% for B unit.



A

B

(28)

20

The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation

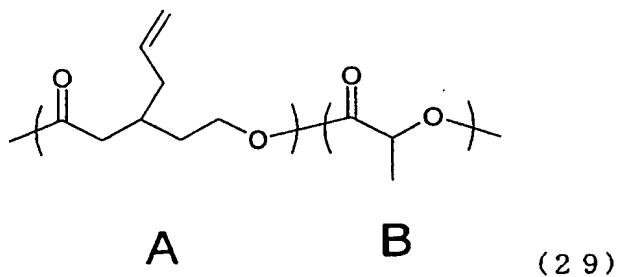
chromatography (GPC; TOSOH HLC-8220, column; TOSOH TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, Mn was 21,500, and weight average 5 molecular weight, Mw was 29,900.

Example 3

(Synthesis of polyester using tetrahydro-4-(2-propenyl)-2H-pyran-2-one and L-lactide)

1.40 g (10.0 mmol) of tetrahydro-4-(2-propenyl)-2H-pyran-2-one represented by the chemical formula (26), 7.21 g (50.0 mmol) of L-lactide, 2.4 ml of 0.1 M toluene solution of tin octylate (tin 2-ethylhexanoate), 2.4 ml of 0.1 M toluene solution of p-tert-butylbenzyl alcohol were placed in a 10 polymerization ampoule and after performing drying under reduced pressure for 1 hour and nitrogen purge, it was heat-sealed under reduced pressure, and heated at 150°C to perform ring-opening polymerization. The 15 reaction was ended after 12 hours and the reaction product was cooled. The obtained polymer was dissolved in chloroform and re-precipitated in 20 methanol in an amount of 10 times of the chloroform required for dissolving the polymer. The precipitation was collected and 6.21 g of polymer was obtained by carrying out drying under reduced 25 pressure. In order to identify the structure of the obtained polymer, NMR analysis was performed on the

same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate copolymer which contains a unit represented by the following chemical formula (29) as a monomer unit. It was also confirmed that the ratios of monomer units were 8 mol% for A unit and 92 mol% for B unit.



The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 23,200, and weight average molecular weight, M_w was 32,200.

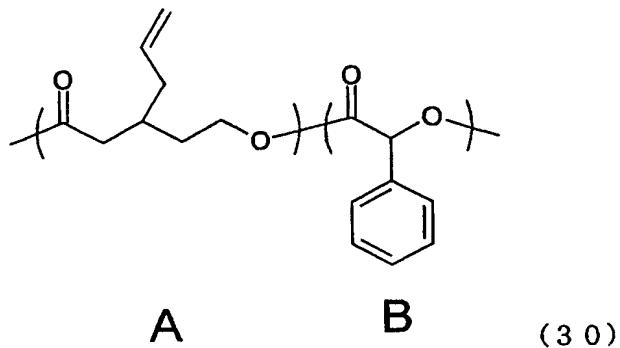
Example 4

(Synthesis of polyester using tetrahydro-4-(2-propenyl)-2H-pyran-2-one and mandelide (3,6-diphenyl-1,4-dioxane-2,5-dione))

20 0.28 g (2.0 mmol) of tetrahydro-4-(2-propenyl)-
2H-pyran-2-one represented by the chemical formula
(26), 2.68 g (10.0 mmol) of mandelide, 4.8 ml of 0.01

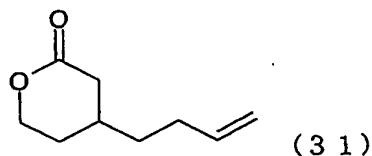
M toluene solution of tin octylate (tin 2-ethylhexanoate), 4.8 ml. of 0.01 M toluene solution of p-tert-butylbenzyl alcohol were placed in a polymerization ampoule and after performing drying under reduced pressure for 1 hour and nitrogen purge, it was heat-sealed under reduced pressure, and heated at 150°C to perform ring-opening polymerization. The reaction was ended after 12 hours and the reaction product was cooled. The obtained polymer was dissolved in chloroform and re-precipitated in methanol in an amount of 10 times of the chloroform required for dissolving the polymer. The precipitation was collected and 1.79 g of polymer was obtained by carrying out drying under reduced pressure.

In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate copolymer which contains a unit represented by the following chemical formula (30) as a monomer unit. It was also confirmed that the ratios of monomer units were 10 mol% for A unit and 90 mol% for B unit.



The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH 5 TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 18,700, and weight average molecular weight, M_w was 28,800.

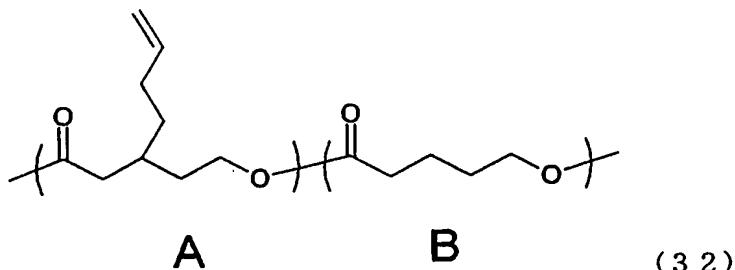
Example 5
 10 (Synthesis of polyester using tetrahydro-4-(2-butenyl)-2H-pyran-2-one represented by chemical formula (31) and δ -valerolactone)



0.28 g (2.0 mmol) of tetrahydro-4-(3-butenyl)-
 15 2H-pyran-2-one represented by the chemical formula (31), 1.00 g (10.0 mmol) of δ -valerolactone, 4.8 ml of 0.01 M toluene solution of tin octylate (tin 2-

ethylhexanoate), 4.8 ml of 0.01 M toluene solution of p-tert-butylbenzyl alcohol were placed in a polymerization ampoule and after performing drying under reduced pressure for 1 hour and nitrogen purge,
5 it was heat-sealed under reduced pressure, and heated at 150°C to perform ring-opening polymerization. The reaction was ended after 12 hours and the reaction product was cooled. The obtained polymer was dissolved in chloroform and re-precipitated in
10 methanol in an amount of 10 times of the chloroform required for dissolving the polymer. The precipitation was collected and 0.89 g of polymer was obtained by carrying out drying under reduced pressure.

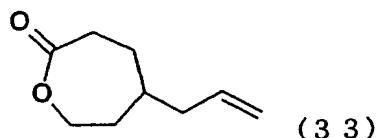
15 In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate copolymer which contains a unit
20 represented by the following chemical formula (32) as a monomer unit. It was also confirmed that the ratios of monomer units were 14 mol% for A unit and 86 mol% for B unit.



The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH 5 TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 21,000, and weight average molecular weight, M_w was 29,000.

Example 6

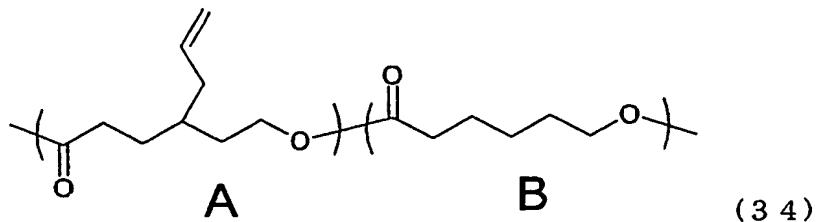
10 (Synthesis of polyester using 5-(2-propenyl)-2-oxepanone represented by chemical formula (33) and ϵ -caprolactone)



15 0.34 g (2.0 mmol) of 5-(2-propenyl)-2-oxepanone represented by the chemical formula (33), 1.14 g (10.0 mmol) of ϵ -caprolactone, 4.8 ml of 0.01 M toluene solution of tin octylate (tin 2-ethylhexanoate), 4.8 ml of 0.01 M toluene solution of

p-tert-butylbenzyl alcohol were placed in a polymerization ampoule and after performing drying under reduced pressure for 1 hour and nitrogen purge, it was heat-sealed under reduced pressure, and heated 5 at 150°C to perform ring-opening polymerization. The reaction was ended after 12 hours and the reaction product was cooled. The obtained polymer was dissolved in chloroform and re-precipitated in methanol in an amount of 10 times of the chloroform 10 required for dissolving the polymer. The precipitation was collected and 0.96 g of polymer was obtained by carrying out drying under reduced pressure.

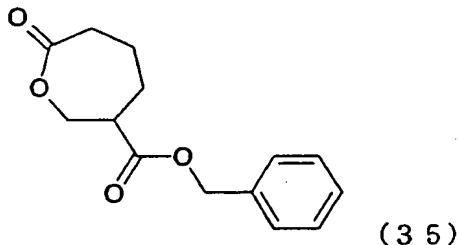
In order to identify the structure of the 15 obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate copolymer which contains a unit represented by the following chemical formula (34) as 20 a monomer unit. It was also confirmed that the ratios of monomer units were 13 mol% for A unit and 87 mol% for B unit.



The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH 5 TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 22,500, and weight average molecular weight, M_w was 32,000.

Example 7

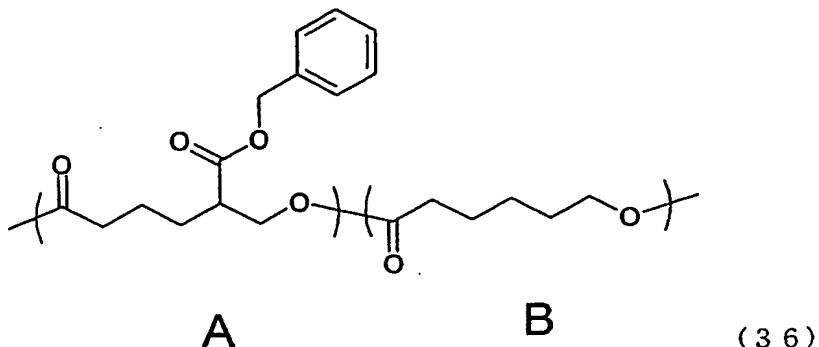
10 (Synthesis of polyester using 7-oxo-3-oxepane
carboxylic acid phenylmethyl ester represented by
chemical formula (35) and ϵ -caprolactone)



0.50 g (2.0 mmol) of 7-oxo-3-oxepane carboxylic acid phenylmethyl ester represented by the chemical formula (35), 1.14 g (10.0 mmol) of ϵ -caprolactone, 4.8 ml of 0.01 M toluene solution of tin octylate

(tin 2-ethylhexanoate), 4.8 ml of 0.01 M toluene solution of p-tert-butylbenzyl alcohol were placed in a polymerization ampoule and after performing drying under reduced pressure for 1 hour and nitrogen purge, 5 it was heat-sealed under reduced pressure, and heated at 150°C to perform ring-opening polymerization. The reaction was ended after 12 hours and the reaction product was cooled. The obtained polymer was dissolved in chloroform and re-precipitated in 10 methanol in an amount of 10 times of the chloroform required for dissolving the polymer. The precipitation was collected and 1.23 g of polymer was obtained by carrying out drying under reduced pressure.

15 In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate copolymer which contains a unit 20 represented by the following chemical formula (36) as a monomer unit. It was also confirmed that the ratios of monomer units were 14 mol% for A unit and 86 mol% for B unit.

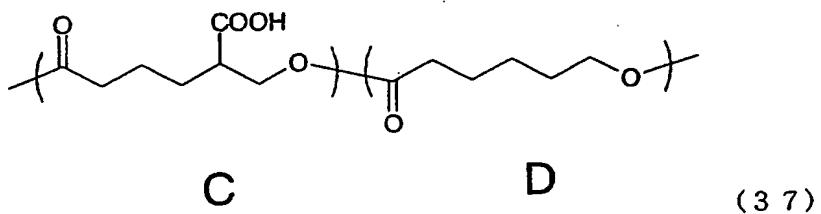


The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH 5 TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 12,000, and weight average molecular weight, M_w was 16,000.

1.00 g of the polyhydroxyalkanoate copolymer
10 represented by the chemical formula (36) obtained
here was dissolved in 100 ml of a mixed solvent of
dioxane-ethanol (75:25), and 0.22 g of 5%
palladium/carbon catalyst was added to this, the
inside of the reaction system was filled with
15 hydrogen, and the reaction mixture was agitated at
room temperature for one day. After the reaction
ended, the reaction mixture was filtered with a 0.25
μm membrane filter in order to remove the catalyst,
and the reaction solution was collected. After the
20 solution was condensed, it was dissolved in

chloroform, re-precipitation was performed using methanol in an amount of 10 times thereof. The obtained polymer was collected and 0.75 g of polymer was obtained by carrying out drying under reduced 5 pressure.

In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a 10 polyhydroxyalkanoate copolymer which contains a unit represented by the following chemical formula (37) as a monomer unit. It was also confirmed that the ratios of monomer units were 14 mol% for C unit and 86 mol% for D unit.

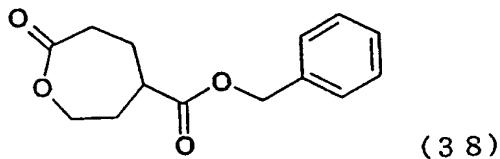


15

The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH TSK-GEL Super HM-H, solvent; chloroform, polystyrene 20 conversion). Consequently, the number average molecular weight, M_n was 10,600, and weight average molecular weight, M_w was 14,700.

Example 8

(Synthesis of polyester using 7-oxo-4-oxepane carboxylic acid phenylmethyl ester represented by chemical formula (38) and L-lactide)



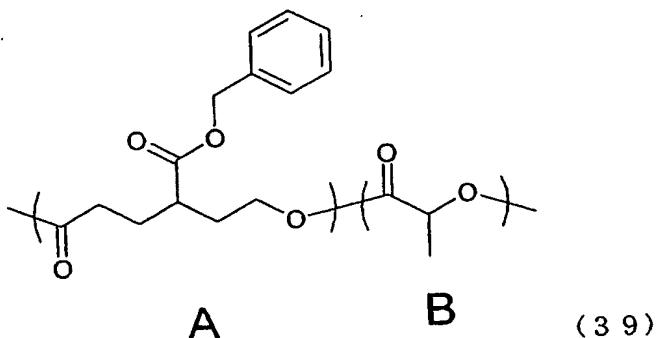
(38)

5

2.48 g (10.0 mmol) of 7-oxo-4-oxepane carboxylic acid phenylmethyl ester represented by the chemical formula (38), 7.21 g (50.0 mmol) of L-lactide, 2.4 ml of 0.1 M toluene solution of tin 10 octylate (tin 2-ethylhexanoate), 2.4 ml of 0.1 M toluene solution of p-tert-butylbenzyl alcohol were placed in a polymerization ampoule and after performing drying under reduced pressure for 1 hour and nitrogen purge, it was heat-sealed under reduced 15 pressure, and heated at 150°C to perform ring-opening polymerization. The reaction was ended after 12 hours and the reaction product was cooled. The obtained polymer was dissolved in chloroform and re-precipitated in methanol in an amount of 10 times of 20 the chloroform required for dissolving the polymer. The precipitation was collected and 7.08 g of polymer was obtained by carrying out drying under reduced pressure.

In order to identify the structure of the

obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate copolymer which contains a unit represented by the following chemical formula (39) as a monomer unit. It was also confirmed that the ratios of monomer units were 8 mol% for A unit and 92 mol% for B unit.

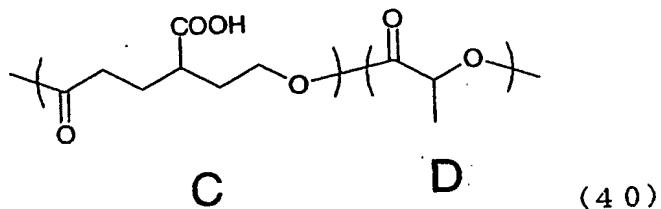


10 The average molecular weight of the obtained
polyhydroxyalkanoate was estimated by gel permeation
chromatography (GPC; TOSOH HLC-8220, column; TOSOH
TSK-GEL Super HM-H, solvent; chloroform, polystyrene
conversion). Consequently, the number average
15 molecular weight, Mn was 10,300, and weight average
molecular weight, Mw was 14,800.

5.00 g of the polyhydroxyalkanoate copolymer represented by the chemical formula (39) obtained here was dissolved in 500 ml of a mixed solvent of 20 dioxane-ethanol (75:25), and 1.10 g of 5% palladium/carbon catalyst was added to this, the

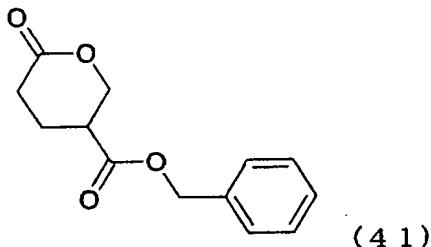
inside of the reaction system was filled with hydrogen, and the reaction mixture was agitated at room temperature for one day. After the reaction ended, the reaction mixture was filtered with a 0.25 5 μm membrane filter in order to remove the catalyst, and the reaction solution was collected. After the solution was condensed, it was dissolved in chloroform, re-precipitation was performed using methanol in an amount of 10 times thereof. The 10 obtained polymer was collected and 3.70 g of polymer was obtained by carrying out drying under reduced pressure.

In order to identify the structure of the obtained polymer, NMR analysis was performed on the 15 same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate copolymer which contains a unit represented by the following chemical formula (40) as a monomer unit. It was also confirmed that the 20 ratios of monomer units were 8 mol% for C unit and 92 mol% for D unit.



The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 9,500, and weight average molecular weight, M_w was 12,900.

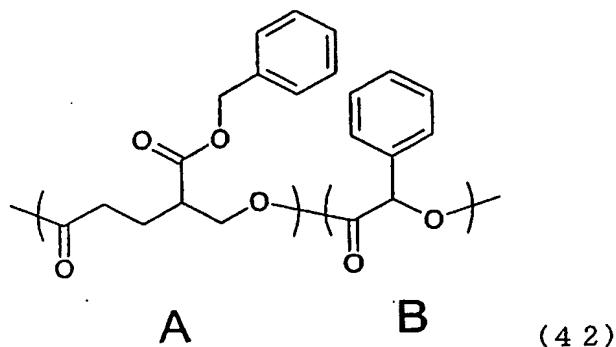
Example 9
(Synthesis of polyester using tetrahydro-6-oxo-2H-pyran-3-carboxylic acid phenylmethyl ester represented by chemical formula (41) and mandelide)



0.47g (2.0 mmol) of tetrahydro-6-oxo-2H-pyran-3-carboxylic acid phenylmethyl ester represented by 15 the chemical formula (41), 2.68 g (10.0 mmol) of mandelide, 4.8 ml of 0.01 M toluene solution of tin octylate (tin 2-ethylhexanoate), 4.8 ml of 0.01 M toluene solution of p-tert-butylbenzyl alcohol were placed in a polymerization ampoule and after 20 performing drying under reduced pressure for 1 hour and nitrogen purge, it was heat-sealed under reduced pressure, and heated at 150°C to perform ring-opening

polymerization. The reaction was ended after 12 hours and the reaction product was cooled. The obtained polymer was dissolved in chloroform and reprecipitated in methanol in an amount of 10 times of the chloroform required for dissolving the polymer. The precipitation was collected and 2.06 g of polymer was obtained by carrying out drying under reduced pressure.

In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate copolymer which contains a unit represented by the following chemical formula (42) as a monomer unit. It was also confirmed that the ratios of monomer units were 7 mol% for A unit and 93 mol% for B unit.



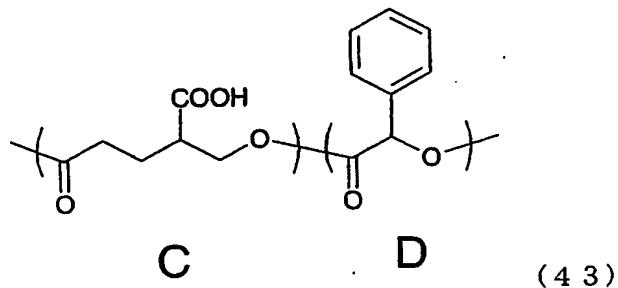
The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation

chromatography (GPC; TOSOH HLC-8220, column; TOSOH TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, Mn was 12,000, and weight average 5 molecular weight, Mw was 16,000.

1.00 g of the polyhydroxyalkanoate copolymer represented by the chemical formula (42) obtained here was dissolved in 100 ml of a mixed solvent of dioxane-ethanol (75:25), and 0.22 g of 5% palladium/carbon catalyst was added to this, the 10 inside of the reaction system was filled with hydrogen, and the reaction mixture was agitated at room temperature for one day. After the reaction ended, the reaction mixture was filtered with a 0.25 15 μm membrane filter in order to remove the catalyst, and the reaction solution was collected. After the solution was condensed, it was dissolved in chloroform, re-precipitation was performed using methanol in an amount of 10 times thereof. The 20 obtained polymer was collected and 0.73 g of polymer was obtained by carrying out drying under reduced pressure.

In order to identify the structure of the obtained polymer, NMR analysis was performed on the 25 same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate copolymer which contains a unit

represented by the following chemical formula (43) as a monomer unit. It was also confirmed that the ratios of monomer units were 7 mol% for C unit and 93 mol% for D unit.



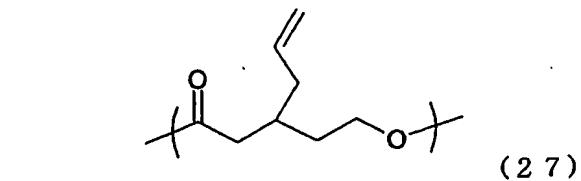
5

The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 8,700, and weight average molecular weight, M_w was 12,900.

10

Example 10

Oxidation reaction of polyhydroxyalkanoate which consists of a unit represented by chemical formula (27) synthesized in Example 1

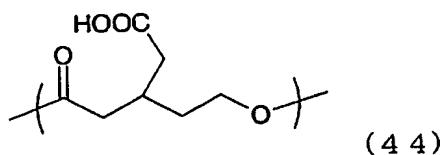


0.50 g of polyhydroxyalkanoate which consists

of a unit represented by the chemical formula (27) obtained in Example 1 was added to an eggplant flask and dissolved with 30 ml of acetone added. This was placed in an ice bath, and 5 ml of acetic acid and 5 2.83 g of 18-crown-6-ether were added and the mixture was agitated. Next, 2.25 g of potassium permanganate was slowly added on the ice bath, and agitated in an ice bath for 2 hours, and agitated at room temperature for further 18 hours. 60 ml of ethyl 10 acetate was added after the reaction ended, and 45 ml of water was further added. Next, sodium hydrogen sulfite was added until peracid was removed. Then, the pH of the liquid was adjusted to 1 with 1.0 N hydrochloric acid. The organic layer was extracted 15 and washed 3 times with 1.0 N hydrochloric acid. After the organic layer was collected, crude polymer was collected by evaporating the solvent. Next, polymer was collected, after washed with 50 ml of water, 50 ml of methanol and further with 50 ml of 20 water 3 times. Next, it was dissolved in 3 ml of THF, re-precipitation was performed using methanol in an amount of 50 times of THF required for dissolving the polymer. The precipitation was collected and 0.42 g of polymer was obtained by carrying out drying under 25 reduced pressure.

In order to identify the structure of the obtained polymer, NMR analysis was performed on the

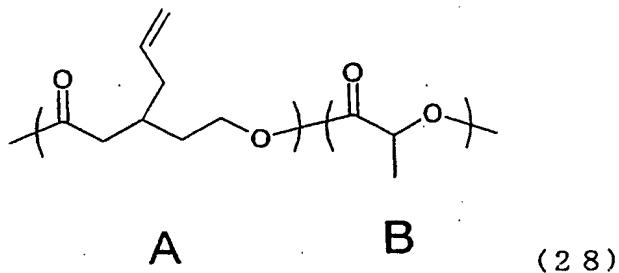
same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate which contains a unit represented by the following chemical formula (44) as a monomer unit.



The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH 10 TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 4,700, and weight average molecular weight, M_w was 6,200.

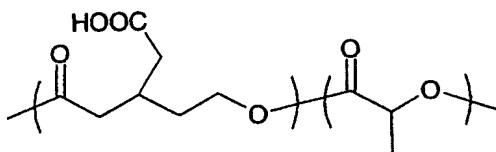
Example 11

15 Oxidation reaction of polyhydroxyalkanoate
which consists of a unit represented by chemical
formula (28) synthesized in Example 2



0.50 g of a polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (28) (A: 7 mol%, B: 93 mol%) obtained in Example 2 was added to an eggplant flask and 5 dissolved with 30 ml of acetone added. This was placed in an ice bath, and 5 ml of acetic acid and 0.36 g of 18-crown-6-ether were added and the mixture was agitated. Next, 0.28 g of potassium permanganate was slowly added on the ice bath, and agitated in an 10 ice bath for 2 hours, and agitated at room temperature for further 18 hours. 60 ml of ethyl acetate was added after the reaction ended, and 45 ml of water was further added. Next, sodium hydrogen sulfite was added until peracid was removed. Then, 15 the pH of the liquid was adjusted to 1 with 1.0 N hydrochloric acid. The organic layer was extracted and washed 3 times with 1.0 N hydrochloric acid. After the organic layer was collected, crude polymer was collected by evaporating the solvent. Next, 20 polymer was collected, after washed with 50 ml of water, 50 ml of methanol and further with 50 ml of water 3 times. Next, it was dissolved in 3 ml of THF, re-precipitation was performed using methanol in an amount of 50 times of THF required for dissolving the 25 polymer. The precipitation was collected and 0.43 g of polymer was obtained by carrying out drying under reduced pressure.

In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a 5 polyhydroxyalkanoate which contains a unit represented by the following chemical formula (45) as a monomer unit.



C

D

(45)

The average molecular weight of the obtained 10 polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 17,400, and weight average 15 molecular weight, M_w was 23,800.

In order to calculate the unit of the obtained polyhydroxyalkanoate furthermore, carboxyl groups at the end of the side chains of the polyhydroxyalkanoate were subjected to methyl 20 esterification using trimethylsilyldiazomethane and the calculation was performed.

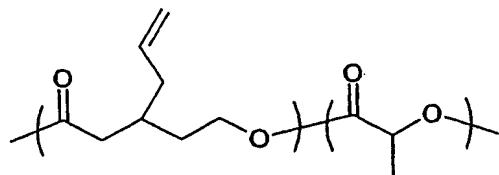
30 mg of the target polyhydroxyalkanoate was

added to 100 ml-volume eggplant flask and dissolved with 2.1 ml of chloroform and 0.7 ml of methanol added. 0.5 ml of 2 mol/L trimethylsilyldiazomethane-hexane solution was added to this, and agitated at 5 room temperature for 1 hour. After the reaction ended, the solvent was removed and the polymer was collected. It was washed with 50 ml of methanol and the polymer was collected after that. 30 mg of polyhydroxyalkanoate was obtained by carrying out 10 drying under reduced pressure.

In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained polyhydroxyalkanoate 15 represented by the chemical formula (45) was a copolymer having a monomer ratio of 7 mol% for C unit and 93 mol% for D unit.

Example 12

Oxidation reaction of polyhydroxyalkanoate 20 which consists of a unit represented by chemical formula (29) synthesized in Example 3



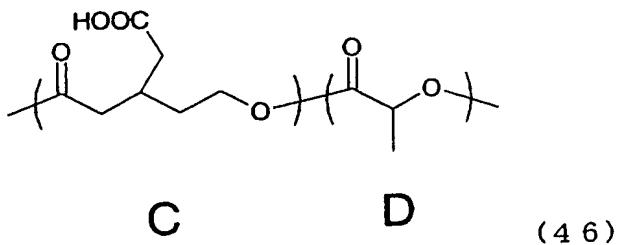
A

B

(29)

5.00 g of a polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (29) (A: 8 mol%, B: 92 mol%) obtained in Example 3 was added to an eggplant flask and 5 dissolved with 300 ml of acetone added. This was placed in an ice bath, and 50 ml of acetic acid and 4.09 g of 18-crown-6-ether were added and the mixture was agitated. Next, 3.26 g of potassium permanganate was slowly added on the ice bath, and agitated in an 10 ice bath for 2 hours, and agitated at room temperature for further 18 hours. 600 ml of ethyl acetate was added after the reaction ended, and 450 ml of water was further added. Next, sodium hydrogen sulfite was added until peracid was removed. Then, 15 the pH of the liquid was adjusted to 1 with 1.0 N hydrochloric acid. The organic layer was extracted and washed 3 times with 1.0 N hydrochloric acid. After the organic layer was collected, crude polymer was collected by evaporating the solvent. Next, 20 polymer was collected, after washed with 500 ml of water, 500 ml of methanol and further with 500 ml of water 3 times. Next, it was dissolved in 30 ml of THF, re-precipitation was performed using methanol in an amount of 50 times of THF required for dissolving 25 the polymer. The precipitation was collected and 4.38 g of polymer was obtained by carrying out drying under reduced pressure.

In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a 5 polyhydroxyalkanoate which contains a unit represented by the following chemical formula (46) as a monomer unit.



C D (46)

The average molecular weight of the obtained 10 polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average 15 molecular weight, Mn was 19,800, and weight average molecular weight, Mw was 27,900.

In order to calculate the unit of the obtained polyhydroxyalkanoate furthermore, carboxyl groups at the end of the side chains of the polyhydroxyalkanoate were subjected to methyl 20 esterification using trimethylsilyldiazomethane and the calculation was performed.

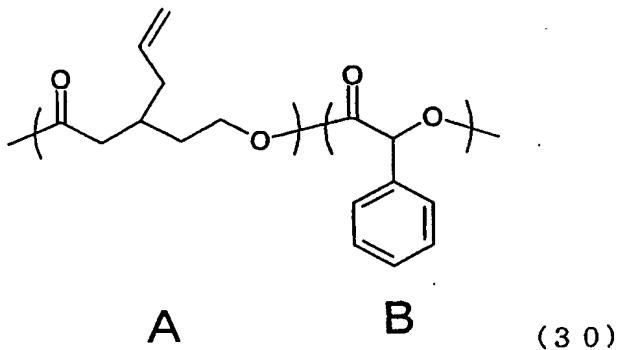
30 mg of the target polyhydroxyalkanoate was

added to 100 ml-volume eggplant flask and dissolved with 2.1 ml of chloroform and 0.7 ml of methanol added. 0.5 ml of 2 mol/L trimethylsilyldiazomethane-hexane solution was added to this, and agitated at 5 room temperature for 1 hour. After the reaction ended, the solvent was removed and the polymer was collected. It was washed with 50 ml of methanol and the polymer was collected after that. 30 mg of polyhydroxyalkanoate was obtained by carrying out 10 drying under reduced pressure.

In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained polyhydroxyalkanoate 15 represented by the chemical formula (46) was a copolymer having a monomer ratio of 8 mol% for C unit and 92 mol% for D unit.

Example 13

Oxidation reaction of polyhydroxyalkanoate 20 which consists of a unit represented by chemical formula (30) synthesized in Example 4



A

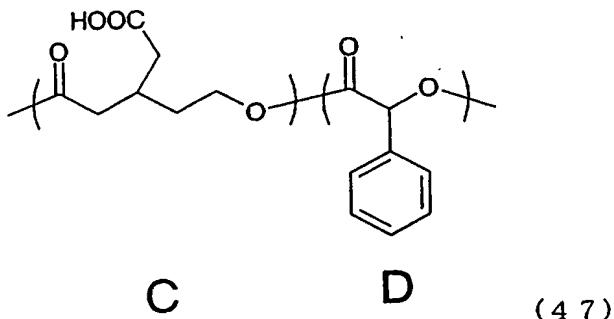
B

(30)

0.50 g of a polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (30) (A: 10 mol%, B: 90 mol%) obtained in 5 Example 4 was added to an eggplant flask and dissolved with 30 ml of acetone added. This was placed in an ice bath, and 5 ml of acetic acid and 0.29 g of 18-crown-6-ether were added and the mixture was agitated. Next, 0.23 g of potassium permanganate 10 was slowly added on the ice bath, and agitated in an ice bath for 2 hours, and agitated at room temperature for further 18 hours. 60 ml of ethyl acetate was added after the reaction ended, and 45 ml of water was further added. Next, sodium hydrogen 15 sulfite was added until peracid was removed. Then, the pH of the liquid was adjusted to 1 with 1.0 N hydrochloric acid. The organic layer was extracted and washed 3 times with 1.0 N hydrochloric acid. After the organic layer was collected, crude polymer 20 was collected by evaporating the solvent. Next,

polymer was collected, after washed with 50 ml of water, 50 ml of methanol and further with 50 ml of water 3 times. Next, it was dissolved in 3 ml of THF, re-precipitation was performed using methanol in an 5 amount of 50 times of THF required for dissolving the polymer. The precipitation was collected and 0.41 g of polymer was obtained by carrying out drying under reduced pressure.

In order to identify the structure of the 10 obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate which contains a unit represented by the following chemical formula (47) as 15 a monomer unit.



The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH 20 TSK-GEL Super HM-H, solvent; chloroform, polystyrene

conversion). Consequently, the number average molecular weight, Mn was 15,500, and weight average molecular weight, Mw was 20,300.

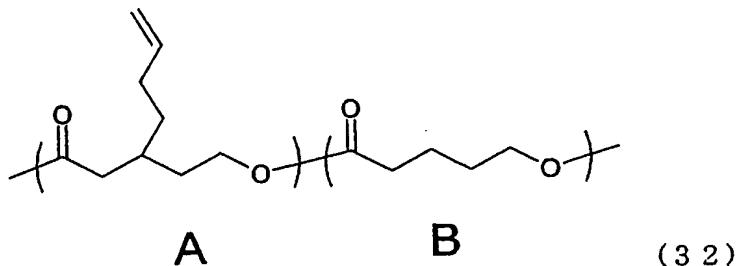
In order to calculate the unit of the obtained 5 polyhydroxyalkanoate furthermore, carboxyl groups at the end of the side chains of the polyhydroxyalkanoate were subjected to methyl esterification using trimethylsilyldiazomethane and the calculation was performed.

10 30 mg of the target polyhydroxyalkanoate was added to 100 ml-volume eggplant flask and dissolved with 2.1 ml of chloroform and 0.7 ml of methanol added. 0.5 ml of 2 mol/L trimethylsilyldiazomethane-hexane solution was added to this, and agitated at 15 room temperature for 1 hour. After the reaction ended, the solvent was removed and the polymer was collected. It was washed with 50 ml of methanol and the polymer was collected after that. 28 mg of polyhydroxyalkanoate was obtained by carrying out 20 drying under reduced pressure.

NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained polyhydroxyalkanoate represented by the chemical formula (47) was a 25 copolymer having a monomer ratio of 10 mol% for C unit and 90 mol% for D unit.

Example 14

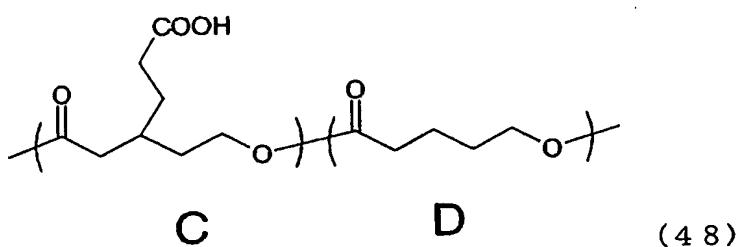
Oxidation reaction of polyhydroxyalkanoate which consists of a unit represented by chemical formula (32) synthesized in Example 5



5 0.50 g of a polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (32) (A: 14 mol%, B: 86 mol%) obtained in Example 5 was added to an eggplant flask and dissolved with 30 ml of acetone added. This was
10 placed in an ice bath, and 5 ml of acetic acid and 0.52 g of 18-crown-6-ether were added and the mixture was agitated. Next, 0.42 g of potassium permanganate was slowly added on the ice bath, and agitated in an ice bath for 2 hours, and agitated at room
15 temperature for further 18 hours. 60 ml of ethyl acetate was added after the reaction ended, and 45 ml of water was further added. Next, sodium hydrogen sulfite was added until peracid was removed. Then, the pH of the liquid was adjusted to 1 with 1.0 N
20 hydrochloric acid. The organic layer was extracted and washed 3 times with 1.0 N hydrochloric acid. After the organic layer was collected, crude polymer

was collected by evaporating the solvent. Next, polymer was collected, after washed with 50 ml of water, 50 ml of methanol and further with 50 ml of water 3 times. Next, it was dissolved in 3 ml of THF, 5 re-precipitation was performed using methanol in an amount of 50 times of THF required for dissolving the polymer. The precipitation was collected and 0.43 g of polymer was obtained by carrying out drying under reduced pressure.

10 In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate which contains a unit 15 represented by the following chemical formula (48) as a monomer unit.



The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation 20 chromatography (GPC; TOSOH HLC-8220, column; TOSOH TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average

molecular weight, Mn was 17,900, and weight average molecular weight, Mw was 24,500.

In order to calculate the unit of the obtained polyhydroxyalkanoate furthermore, carboxyl groups at 5 the end of the side chains of the polyhydroxyalkanoate were subjected to methyl esterification using trimethylsilyldiazomethane and the calculation was performed.

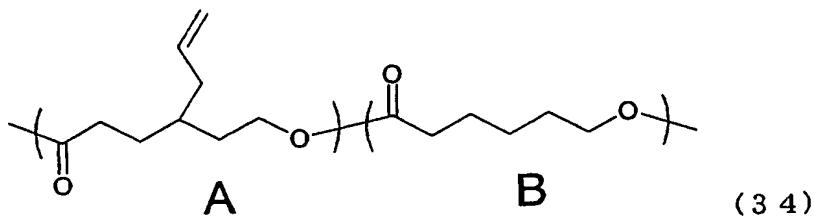
30 mg of the target polyhydroxyalkanoate was 10 added to 100 ml-volume eggplant flask and dissolved with 2.1 ml of chloroform and 0.7 ml of methanol added. 0.5 ml of 2 mol/L trimethylsilyldiazomethane-hexane solution was added to this, and agitated at room temperature for 1 hour. After the reaction 15 ended, the solvent was removed and the polymer was collected. It was washed with 50 ml of methanol and the polymer was collected after that. 29 mg of polyhydroxyalkanoate was obtained by carrying out drying under reduced pressure.

20 NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained polyhydroxyalkanoate represented by the chemical formula (48) was a copolymer having a monomer ratio of 13 mol% for C 25 unit and 87 mol% for D unit.

Example 15

Oxidation reaction of polyhydroxyalkanoate

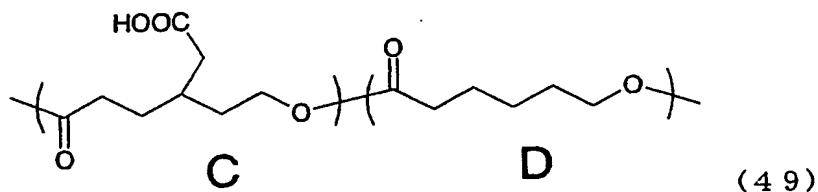
which consists of a unit represented by chemical formula (34) synthesized in Example 6



0.50 g of a polyhydroxyalkanoate copolymer
5 which consists of a unit represented by the chemical formula (34) (A: 13 mol%, B: 87 mol%) obtained in Example 6 was added to an eggplant flask and dissolved with 30 ml of acetone added. This was placed in an ice bath, and 5 ml of acetic acid and
10 0.43 g of 18-crown-6-ether were added and the mixture was agitated. Next, 0.34 g of potassium permanganate was slowly added on the ice bath, and agitated in an ice bath for 2 hours, and agitated at room temperature for further 18 hours. 60 ml of ethyl
15 acetate was added after the reaction ended, and 45 ml of water was further added. Next, sodium hydrogen sulfite was added until peracid was removed. Then, the pH of the liquid was adjusted to 1 with 1.0 N hydrochloric acid. The organic layer was extracted
20 and washed 3 times with 1.0 N hydrochloric acid. After the organic layer was collected, crude polymer was collected by evaporating the solvent. Next,

polymer was collected, after washed with 50 ml of water, 50 ml of methanol and further with 50 ml of water 3 times. Next, it was dissolved in 3 ml of THF, re-precipitation was performed using methanol in an 5 amount of 50 times of THF required for dissolving the polymer. The precipitation was collected and 0.44 g of polymer was obtained by carrying out drying under reduced pressure.

In order to identify the structure of the 10 obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate which contains a unit represented by the following chemical formula (49) as 15 a monomer unit.



The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH 20 TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 18,900, and weight average

molecular weight, Mw was 27,600.

In order to calculate the unit of the obtained polyhydroxyalkanoate furthermore, carboxyl groups at the end of the side chains of the 5 polyhydroxyalkanoate were subjected to methyl esterification using trimethylsilyldiazomethane and the calculation was performed.

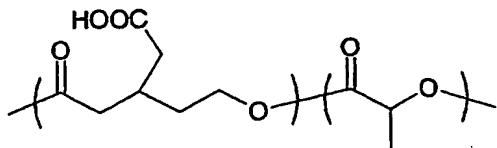
30 mg of the target polyhydroxyalkanoate was added to 100 ml-volume eggplant flask and dissolved 10 with 2.1 ml of chloroform and 0.7 ml of methanol added. 0.5 ml of 2 mol/L trimethylsilyldiazomethane-hexane solution was added to this, and agitated at room temperature for 1 hour. After the reaction ended, the solvent was removed and the polymer was 15 collected. It was washed with 50 ml of methanol and the polymer was collected after that. 28 mg of polyhydroxyalkanoate was obtained by carrying out drying under reduced pressure.

NMR analysis was performed on the same 20 conditions as in Example 1 and consequently it was confirmed that the obtained polyhydroxyalkanoate represented by the chemical formula (49) was a copolymer having a monomer ratio of 12 mol% for C unit and 88 mol% for D unit.

25 Example 16

Condensation reaction of polyhydroxyalkanoate which consist of a unit represented by chemical

formula (46) synthesized in Example 12 and 2-
aminobenzenesulfonic acid



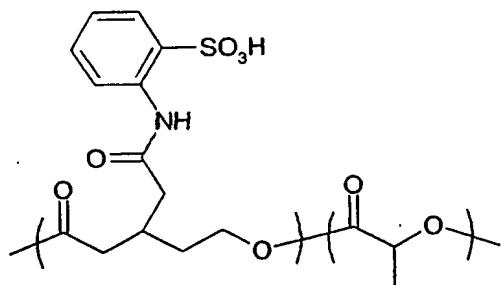
C D
(46)

In a nitrogen atmosphere, 0.40 g of the
5 polyhydroxyalkanoate copolymer which consists of a
unit represented by the chemical formula (46) (C: 8
mol%, D: 92 mol%) obtained in Example 12, 0.35 g of
2-aminobenzenesulfonic acid were added to a 100 ml
three-necked flask and agitated with 15.0 ml of
10 pyridine added, and then 1.06 ml of triphenyl
phosphite was added and heated at 120°C for 6 hours.
After the reaction ended, re-precipitation was
performed from 150 ml of ethanol to collect the
product. The obtained polymer was washed with 1N
15 hydrochloric acid for one day and further washed by
agitating in water for one day, and 0.34 g of polymer
was obtained by carrying out drying under reduced
pressure.

The structure of the obtained polymer was
20 determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker
DPX400; resonant frequency: 400 MHz; nuclide
measured: ^1H ; solvent used: d-DMSO; measurement

temperature: room temperature) and Fourier transformation-infrared absorption (FT-IR) spectrum (Nicolet AVATAR360 FT-IR). As a result of performing IR measurement, the peak of 1,695 cm⁻¹ resulted from 5 carboxylic acid decreased, and a peak resulted from amide group at 1,658 cm⁻¹ was newly observed.

The results of ¹H-NMR showed a shift of the peak resulted from the aromatic ring in 2-aminobenzenesulfonic acid structure and accordingly 10 it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (50) as a monomer unit.



E

F

(50)

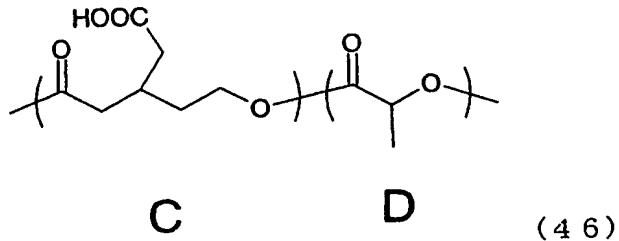
As for the ratio of the units in the 15 polyhydroxyalkanoate represented by the chemical formula (50), it was confirmed that it was a copolymer having a monomer ratio of 8 mol% for unit E and 92 mol % for unit F.

The average molecular weight of the obtained 20 polyhydroxyalkanoate was estimated by gel permeation

chromatography (GPC; TOSOH HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, Mn was 16,700, 5 and weight average molecular weight, Mw was 24,700.

Example 17

Condensation reaction of polyhydroxyalkanoate which consist of a unit represented by chemical formula (46) synthesized in Example 12 and 4-10 aminobenzenesulfonic acid

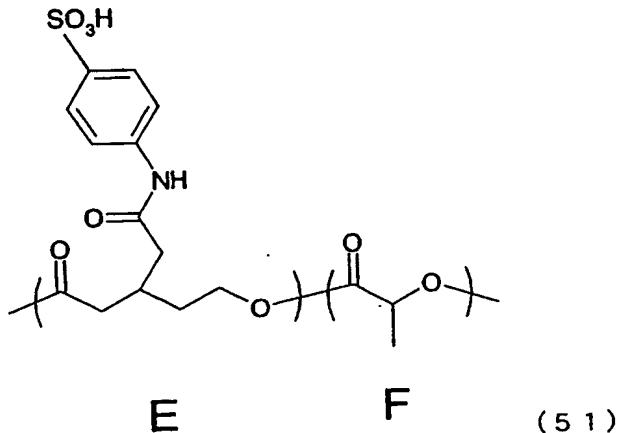


In a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (46) (C: 8 15 mol%, D: 92 mol%) obtained in Example 12, 0.35 g of 4-aminobenzenesulfonic acid were added to a 100 ml three-necked flask and agitated with 15.0 ml of pyridine added, and then 1.06 ml of triphenyl phosphite was added and heated at 120°C for 6 hours. 20 After the reaction ended, re-precipitation was performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N

hydrochloric acid for one day and further washed by agitating in water for one day, and 0.32 g of polymer was obtained by carrying out drying under reduced pressure.

5 The structure of the obtained polymer was determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier 10 transformation-infrared absorption (FT-IR) spectrum (Nicolet AVATAR360 FT-IR). As a result of performing IR measurement, the peak of $1,695\text{ cm}^{-1}$ resulted from carboxylic acid decreased, and a peak resulted from amide group at $1,658\text{ cm}^{-1}$ was newly observed.

15 The results of $^1\text{H-NMR}$ showed a shift of the peak resulted from the aromatic ring in 4-aminobenzenesulfonic acid structure and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by 20 the following chemical formula (51) as a monomer unit.

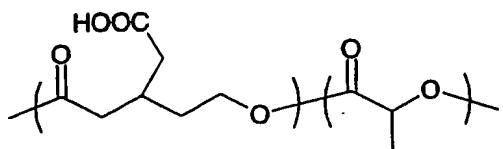


As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (51), it was confirmed that it was a 5 copolymer having a monomer ratio of 8 mol% for unit E and 92 mol % for unit F.

The average molecular weight of the obtained polymer was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer 10 Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 16,100, and weight average molecular weight, M_w was 24,300.

Example 18
 15 Condensation reaction of the polyhydroxyalkanoate which consist of a unit represented by chemical formula (46) synthesized in Example 12 and p-toluidine-2-sulfonic acid

100



C

D

(46)

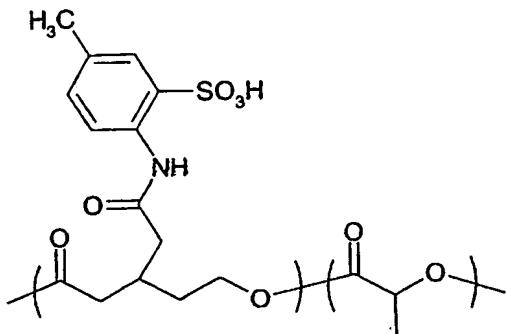
In a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (46) (C: 8 mol%, D: 92 mol%) obtained in Example 12, 0.38 g of p-toluidine-2-sulfonic acid were added to a 100 ml three-necked flask and agitated with 15.0 ml of pyridine added, and then 1.06 ml of triphenyl phosphite was added and heated at 120°C for 6 hours.

After the reaction ended, re-precipitation was performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N hydrochloric acid for one day and further washed by agitating in water for one day, and 0.35 g of polymer was obtained by carrying out drying under reduced pressure.

The structure of the obtained polymer was determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier transformation-infrared absorption (FT-IR) spectrum

(Nicolet AVATAR360 FT-IR). As a result of performing IR measurement, the peak of 1,695 cm⁻¹ resulted from carboxylic acid decreased, and a peak resulted from amide group at 1,658 cm⁻¹ was newly observed.

5 The results of ¹H-NMR showed a shift of the peak resulted from the aromatic ring in p-toluidine-2-sulfonic acid structure and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by
10 the following chemical formula (52) as a monomer unit.



E F (52)

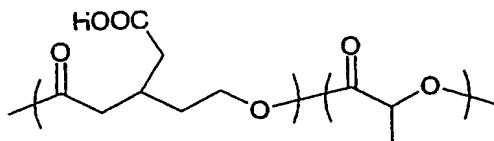
As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (52), it was confirmed that it was a
15 copolymer having a monomer ratio of 8 mol% for unit E and 92 mol % for unit F.

The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer

Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, Mn was 15,700, and weight average molecular weight, Mw was 24,600.

5 Example 19

Condensation reaction of polyhydroxyalkanoate which consist of a unit represented by chemical formula (46) synthesized in Example 12 and 2-aminobenzenesulfonic acid phenyl ester



C

D

(46)

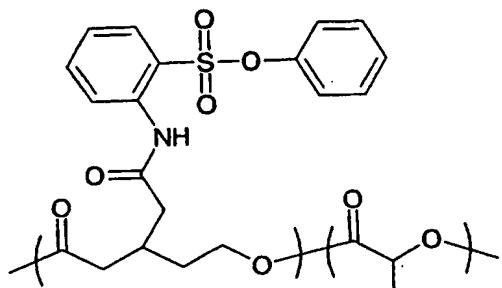
10

In a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (46) (C: 8 mol%, D: 92 mol%) obtained in Example 12, 0.51 g of 15 2-aminobenzenesulfonic acid phenyl ester were added to a 100 ml three-necked flask and agitated with 15.0 ml of pyridine added, and then 1.06 ml of triphenyl phosphite was added and heated at 120°C for 6 hours. After the reaction ended, re-precipitation was 20 performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N hydrochloric acid for one day and further washed by

agitating in water for one day, and 0.38 g of polymer was obtained by carrying out drying under reduced pressure.

The structure of the obtained polymer was 5 determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier transformation-infrared absorption (FT-IR) spectrum 10 (Nicolet AVATAR360 FT-IR). As a result of performing IR measurement, the peak of $1,695\text{ cm}^{-1}$ resulted from carboxylic acid decreased, and a peak resulted from amide group at $1,658\text{ cm}^{-1}$ was newly observed.

The results of $^1\text{H-NMR}$ showed a shift of the peak 15 resulted from the aromatic ring in 2-aminobenzenesulfonic acid phenyl ester structure and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (53) as 20 a monomer unit.



E

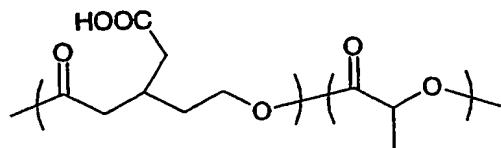
F

(53)

As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (53), it was confirmed that it was a 5 copolymer having a monomer ratio of 8 mol% for unit E and 92 mol % for unit F.

The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer 10 Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 17,100, and weight average molecular weight, M_w was 24,600.

Example 20
 15 Condensation reaction of the polyhydroxyalkanoate which consist of a unit represented by chemical formula (46) synthesized in Example 12 and 2-amino-1-naphthalenesulfonic acid



C

D

(46)

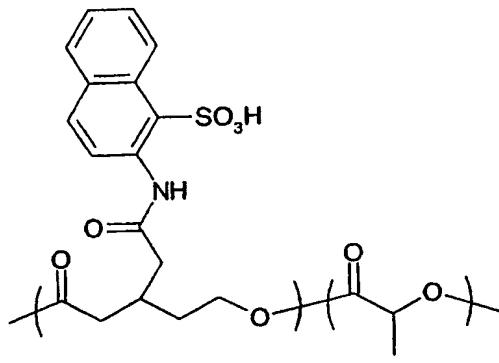
In a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (46) (C: 8 mol%, D: 92 mol%) obtained in Example 12, 0.45 g of 2-amino-1-naphthalenesulfonic acid were added to a 100 ml three-necked flask and agitated with 15.0 ml of pyridine added, and then 1.06 ml of triphenyl phosphite was added and heated at 120°C for 6 hours.

After the reaction ended, re-precipitation was performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N hydrochloric acid for one day and further washed by agitating in water for one day, and 0.34 g of polymer was obtained by carrying out drying under reduced pressure.

The structure of the obtained polymer was determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier transformation-infrared absorption (FT-IR) spectrum

(Nicolet AVATAR360 FT-IR). As a result of performing IR measurement, the peak of 1,695 cm⁻¹ resulted from carboxylic acid decreased, and a peak resulted from amide group at 1,658 cm⁻¹ was newly observed.

5 The results of ¹H-NMR showed a shift of the peak resulted from the aromatic ring in 2-amino-1-naphthalenesulfonic acid structure and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by
10 10 the following chemical formula (54) as a monomer unit.



E F (54)

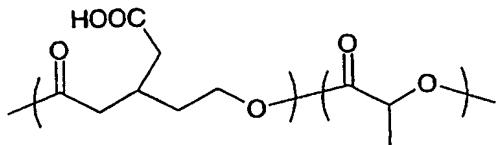
As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (54), it was confirmed that it was a
15 copolymer having a monomer ratio of 8 mol% for unit E and 92 mol % for unit F.

The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer

Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, Mn was 16,900, and weight average molecular weight, Mw was 25,700.

5 Example 21

Condensation reaction of the polyhydroxyalkanoate which consist of a unit represented by chemical formula (46) synthesized in Example 12 and 2-amino-2-methylpropanesulfonic acid



C

D

(46)

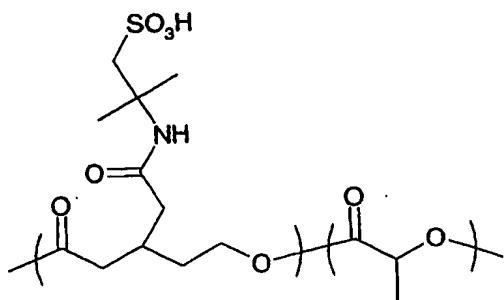
10

In a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (46) (C: 8 mol%, D: 92 mol%) obtained in Example 12, 0.31 g of 15 2-amino-2-methylpropanesulfonic acid were added to a 100 ml three-necked flask and agitated with 15.0 ml of pyridine added, and then 1.06 ml of triphenyl phosphite was added and heated at 120°C for 6 hours. After the reaction ended, re-precipitation was 20 performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N hydrochloric acid for one day and further washed by

agitating in water for one day, and 0.31 g of polymer was obtained by carrying out drying under reduced pressure.

The structure of the obtained polymer was determined by analyzing with ^1H -NMR (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier transformation-infrared absorption (FT-IR) spectrum (Nicolet AVATAR360 FT-IR). As a result of performing IR measurement, the peak of $1,695\text{ cm}^{-1}$ resulted from carboxylic acid decreased, and a peak resulted from amide group at $1,668\text{ cm}^{-1}$ was newly observed.

The results of $^1\text{H-NMR}$ showed a shift of the peak
15 resulted from the methylene of 2-amino-2-methylpropanesulfonic acid structure and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (55) as a monomer unit.



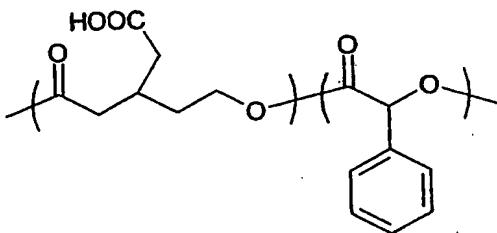
E F (55)

As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (55), it was confirmed that it was a copolymer having a monomer ratio of 8 mol% for unit E 5 and 92 mol% for unit F.

The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 10 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 17,300, and weight average molecular weight, M_w was 24,600.

Example 22

Condensation reaction of polyhydroxyalkanoate 15 which consist of a unit represented by chemical formula (47) synthesized in Example 13 and 3-aminobenzenesulfonic acid



C

D

(47)

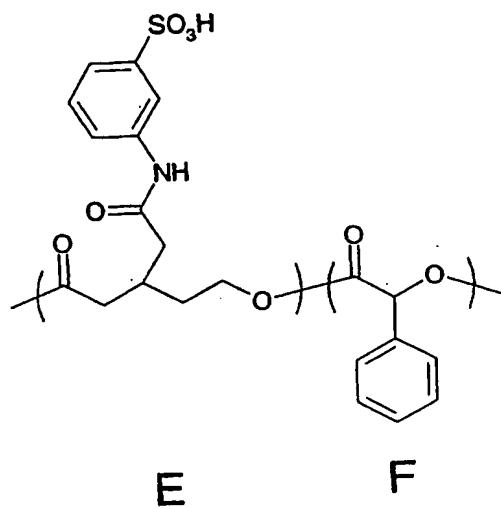
In a nitrogen atmosphere, 0.40 g of the 20 polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (47) (C: 10

mol%, D: 90 mol%) obtained in Example 13, 0.25 g of 3-aminobenzenesulfonic acid were added to a 100 ml three-necked flask and agitated with 15.0 ml of pyridine added, and then 0.76 ml of triphenyl 5 phosphite was added and heated at 120°C for 6 hours. After the reaction ended, re-precipitation was performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N hydrochloric acid for one day and further washed by 10 agitating in water for one day, and 0.33 g of polymer was obtained by carrying out drying under reduced pressure.

The structure of the obtained polymer was determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker 15 DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier transformation-infrared absorption (FT-IR) spectrum (Nicolet AVATAR360 FT-IR). As a result of performing 20 IR measurement, the peak of $1,695\text{ cm}^{-1}$ resulted from carboxylic acid decreased, and a peak resulted from amide group at $1,658\text{ cm}^{-1}$ was newly observed.

The results of $^1\text{H-NMR}$ showed a shift of the peak resulted from the aromatic ring of 3- 25 aminobenzenesulfonic acid structure and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by

the following chemical formula (56) as a monomer unit.



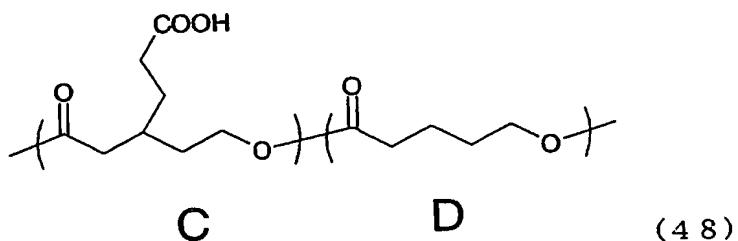
As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (56), it was confirmed that it was a 5 copolymer having a monomer ratio of 10 mol% for unit E and 90 mol% for unit F.

The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation 10 chromatography (GPC; TOSOH HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 13,100, and weight average molecular weight, M_w was 17,700.

15 Example 23

Condensation reaction of the polyhydroxyalkanoate which consist of a unit represented by chemical formula (48) synthesized in

Example 14 and 4-methoxyaniline-2-sulfonic acid

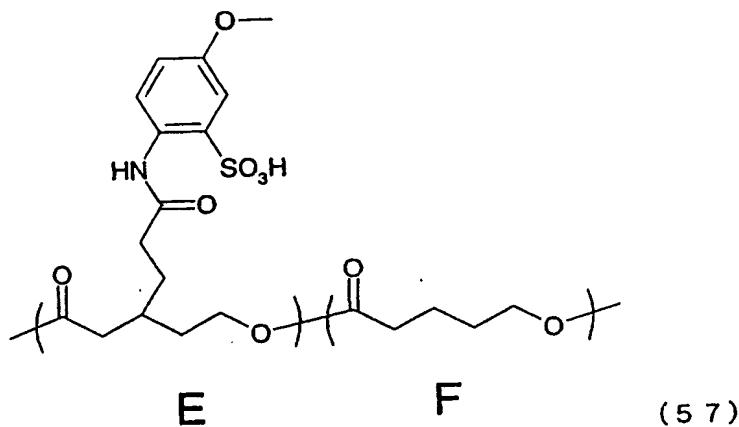


In a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer which consists of a 5 unit represented by the chemical formula (48) (C: 13 mol%, D: 87 mol%) obtained in Example 14, 0.48g of 4-methoxyaniline-2-sulfonic acid were added to a 100 ml three-necked flask and agitated with 15.0 ml of pyridine added, and then 1.24 ml of triphenyl 10 phosphite was added and heated at 120°C for 6 hours. After the reaction ended, re-precipitation was performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N hydrochloric acid for one day and further washed by 15 agitating in water for one day, and 0.37 g of polymer was obtained by carrying out drying under reduced pressure.

The structure of the obtained polymer was determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker 20 DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier

transformation-infrared absorption (FT-IR) spectrum (Nicolet AVATAR360 FT-IR). As a result of performing IR measurement, the peak of $1,695\text{ cm}^{-1}$ resulted from carboxylic acid decreased, and a peak resulted from 5 amide group at $1,658\text{ cm}^{-1}$ was newly observed.

The results of $^1\text{H-NMR}$ showed a shift of the peak resulted from the aromatic ring of 4-methoxyaniline-2-sulfonic acid structure and accordingly it was confirmed that the obtained polymer was a 10 polyhydroxyalkanoate containing a unit represented by the following chemical formula (57) as a monomer unit.



As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical 15 formula (57), it was confirmed that it was a copolymer having a monomer ratio of 13 mol% for unit E and 87 mol% for unit F.

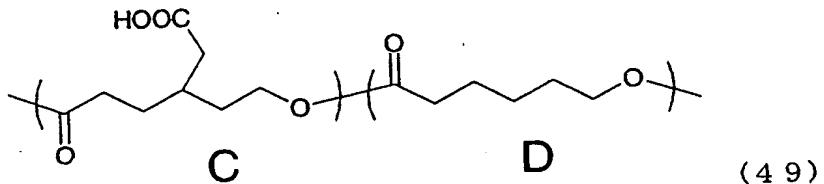
The average molecular weight of the obtained polymer was estimated by gel permeation

chromatography (GPC; TOSOH HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 14,800, 5 and weight average molecular weight, M_w was 20,600.

Example 24

Condensation reaction of the polyhydroxyalkanoate which consist of a unit represented by chemical formula (49) synthesized in

10 Example 15 and taurine

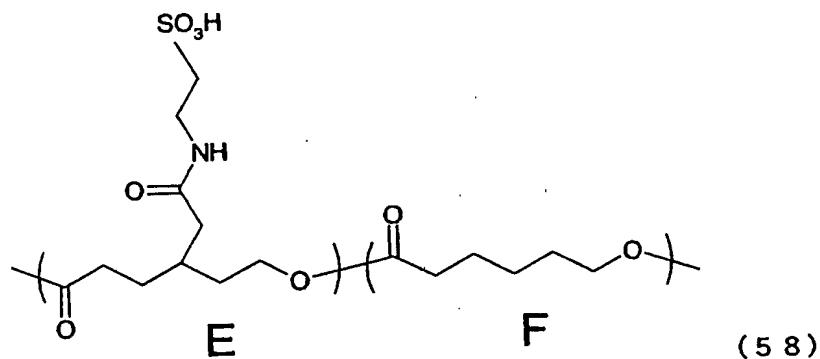


In a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (49) (C: 12 15 mol%, D: 88 mol%) obtained in Example 15, 0.25 g of taurine were added to a 100 ml three-necked flask and agitated with 15.0 ml of pyridine added, and then 1.03 ml of triphenyl phosphite was added and heated at 120°C for 6 hours. After the reaction ended, re- 20 precipitation was performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N hydrochloric acid for one day and further washed by agitating in water for one day, and 0.31 g

of polymer was obtained by carrying out drying under reduced pressure.

The structure of the obtained polymer was determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker 5 DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier transformation-infrared absorption (FT-IR) spectrum (Nicolet AVATAR360 FT-IR). As a result of performing 10 IR measurement, the peak of $1,695\text{ cm}^{-1}$ resulted from carboxylic acid decreased, and a peak resulted from amide group at $1,668\text{ cm}^{-1}$ was newly observed.

The results of $^1\text{H-NMR}$ showed a shift of the peak resulted from the methylene of taurine structure and 15 accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (58) as a monomer unit.



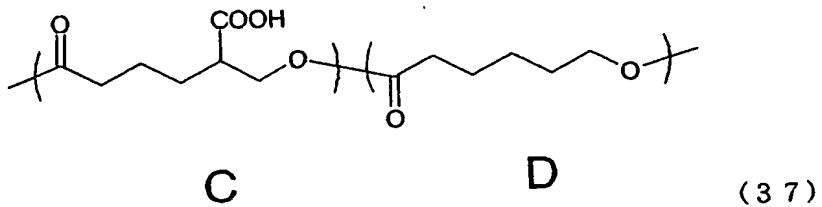
20 As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical

formula (58), it was confirmed that it was a copolymer having a monomer ratio of 12 mol% for unit E and 88 mol% for unit F.

The average molecular weight of the obtained polymer was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 15,500, and weight average molecular weight, M_w was 22,200.

Example 25

Condensation reaction of polyhydroxyalkanoate which consist of a unit represented by chemical formula (37) synthesized in Example 7 and 2-15 aminobenzenesulfonic acid

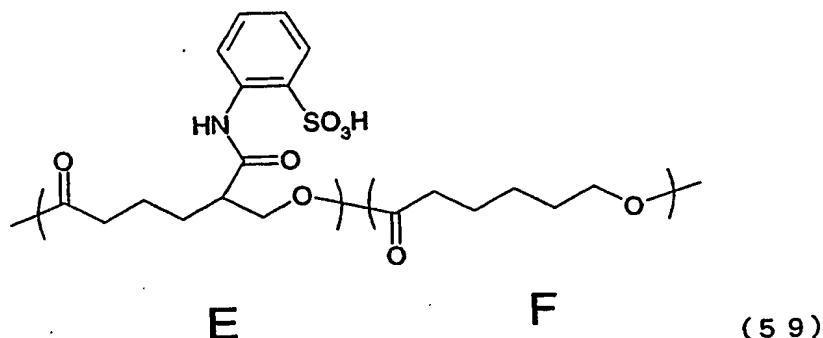


In a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (37) (C: 14 mol%, D: 86 mol%) obtained in Example 7, 0.40 g of 2-aminobenzenesulfonic acid were added to a 100 ml three-necked flask and agitated with 15.0 ml of

pyridine added, and then 1.21 ml of triphenyl phosphite was added and heated at 120°C for 6 hours. After the reaction ended, re-precipitation was performed from 150 ml of ethanol to collect the 5 product. The obtained polymer was washed with 1N hydrochloric acid for one day and further washed by agitating in water for one day, and 0.36 g of polymer was obtained by carrying out drying under reduced pressure.

10 The structure of the obtained polymer was determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier 15 transformation-infrared absorption (FT-IR) spectrum (Nicolet AVATAR360 FT-IR). As a result of performing IR measurement, the peak of $1,695\text{ cm}^{-1}$ resulted from carboxylic acid decreased, and a peak resulted from amide group at $1,658\text{ cm}^{-1}$ was newly observed.

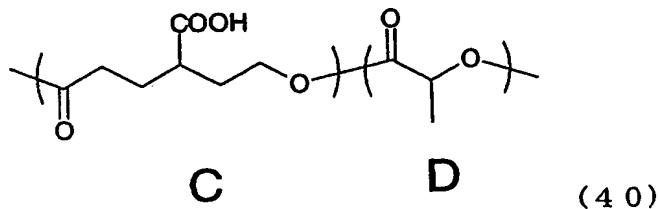
20 The results of $^1\text{H-NMR}$ showed a shift of the peak resulted from the aromatic ring of 2-aminobenzenesulfonic acid structure and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by 25 the following chemical formula (59) as a monomer unit.



As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (59), it was confirmed that it was a 5 copolymer having a monomer ratio of 14 mol% for unit E and 86 mol% for unit F.

The average molecular weight of the obtained polymer was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer 10 Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 9,400, and weight average molecular weight, M_w was 13,400.

Example 26
 15 Condensation reaction of the polyhydroxyalkanoate which consist of a unit represented by chemical formula (40) synthesized in Example 8 and 2-amino-2-methylpropanesulfonic acid

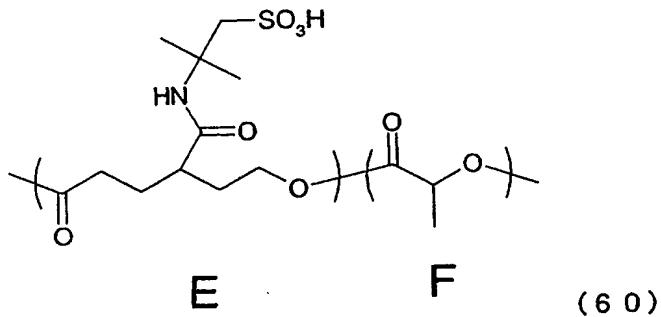


In a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (40) (C: 8 mol%, D: 92 mol%) obtained in Example 8, 0.31 g of 2-amino-2-methylpropanesulfonic acid were added to a 100 ml three-necked flask and agitated with 15.0 ml of pyridine added, and then 1.06 ml of triphenyl phosphite was added and heated at 120°C for 6 hours. After the reaction ended, re-precipitation was performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N hydrochloric acid for one day and further washed by agitating in water for one day, and 0.33 g of polymer was obtained by carrying out drying under reduced pressure.

The structure of the obtained polymer was determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier transformation-infrared absorption (FT-IR) spectrum (Nicolet AVATAR360 FT-IR). As a result of performing

IR measurement, the peak of $1,695\text{ cm}^{-1}$ resulted from carboxylic acid decreased, and a peak resulted from amide group at $1,668\text{ cm}^{-1}$ was newly observed.

The results of $^1\text{H-NMR}$ showed a shift of the peak 5 resulted from the methylene of 2-amino-2-methylpropanesulfonic acid structure and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (60) as a monomer unit.



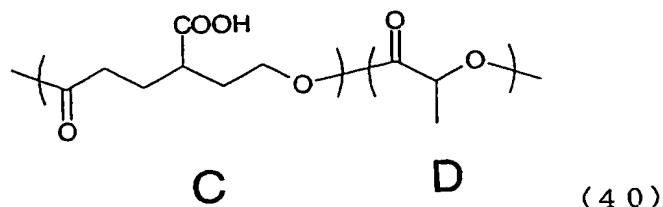
As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (60), it was confirmed that it was a copolymer having a monomer ratio of 8 mol% for unit E 15 and 92 mol% for unit F.

The average molecular weight of the obtained polymer was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 20 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 8,100,

and weight average molecular weight, Mw was 12,400.

Example 27

Condensation reaction of polyhydroxyalkanoate which consist of a unit represented by chemical formula (40) synthesized in Example 8 and 4-aminobenzenesulfonic acid phenyl ester

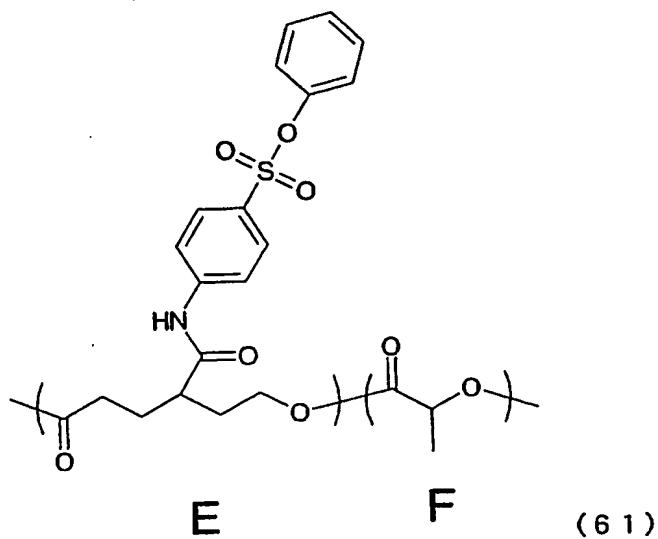


In a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer which consists of a 10 unit represented by the chemical formula (40) (C: 8 mol%, D: 92 mol%) obtained in Example 8, 0.51 g of 4-aminobenzenesulfonic acid phenyl ester were added to a 100 ml three-necked flask and agitated with 15.0 ml of pyridine added, and then 1.06 ml of triphenyl phosphite was added and heated at 120°C for 6 hours. After the reaction ended, re-precipitation was performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N hydrochloric acid for one day and further washed by 15 agitating in water for one day, and 0.35 g of polymer was obtained by carrying out drying under reduced 20 pressure.

The structure of the obtained polymer was

determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier transformation-infrared absorption (FT-IR) spectrum (Nicolet AVATAR360 FT-IR). As a result of performing IR measurement, the peak of $1,695\text{ cm}^{-1}$ resulted from carboxylic acid decreased, and a peak resulted from amide group at $1,658\text{ cm}^{-1}$ was newly observed.

The results of $^1\text{H-NMR}$ showed a shift of the peak resulted from the aromatic ring of 4-aminobenzenesulfonic acid phenyl ester structure and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (61) as a monomer unit.

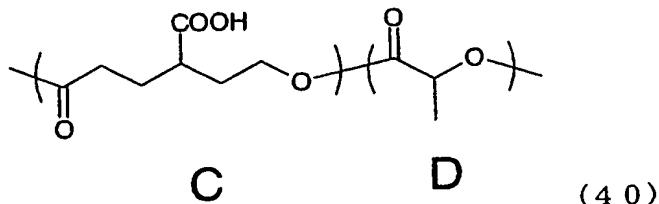


As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (61), it was confirmed that it was a copolymer having a monomer ratio of 8 mol% for unit E 5 and 92 mol% for unit F.

The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 10 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 7,800, and weight average molecular weight, M_w was 14,300.

Example 28

Condensation reaction of the 15 polyhydroxyalkanoate which consist of a unit represented by chemical formula (40) synthesized in Example 8 and 1-naphthylamine-8-sulfonic acid

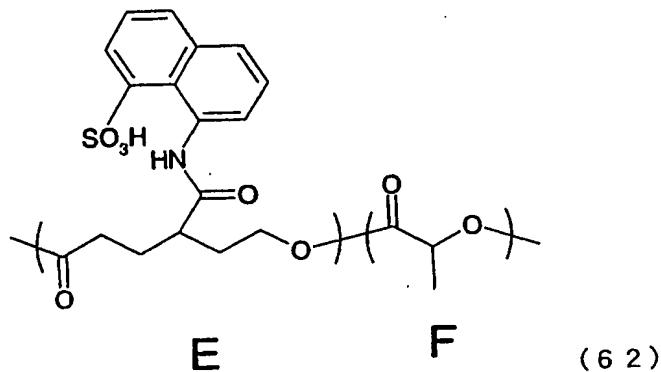


In a nitrogen atmosphere, 0.40 g of the 20 polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (40) (C: 8 mol%, D: 92 mol%) obtained in Example 8, 0.45 g of 1-naphthylamine-8-sulfonic acid were added to a 100 ml

three-necked flask and agitated with 15.0 ml of pyridine added, and then 1.06 ml of triphenyl phosphite was added and heated at 120°C for 6 hours. After the reaction ended, re-precipitation was 5 performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N hydrochloric acid for one day and further washed by agitating in water for one day, and 0.33 g of polymer was obtained by carrying out drying under reduced 10 pressure.

The structure of the obtained polymer was determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement 15 temperature: room temperature) and Fourier transformation-infrared absorption (FT-IR) spectrum (Nicolet AVATAR360 FT-IR). As a result of performing IR measurement, the peak of $1,695\text{ cm}^{-1}$ resulted from carboxylic acid decreased, and a peak resulted from 20 amide group at $1,658\text{ cm}^{-1}$ was newly observed.

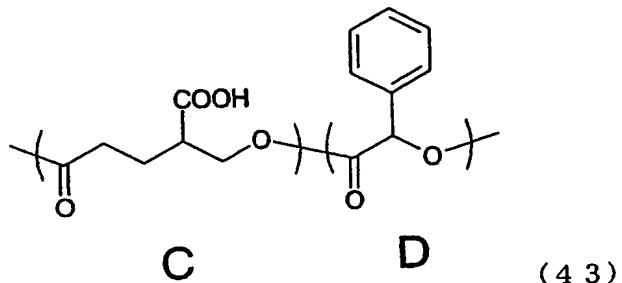
The results of $^1\text{H-NMR}$ showed a shift of the peak resulted from the aromatic ring of 1-naphthylamine-8-sulfonic acid structure and accordingly it was confirmed that the obtained polymer was a 25 polyhydroxyalkanoate containing a unit represented by the following chemical formula (62) as a monomer unit.



As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (62), it was confirmed that it was a 5 copolymer having a monomer ratio of 8 mol% for unit E and 92 mol% for unit F.

The average molecular weight of the obtained polymer was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer 10 Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, Mn was 8,200, and weight average molecular weight, Mw was 12,400.

Example 29
15 Condensation reaction of the polyhydroxyalkanoate which consist of a unit represented by chemical formula (43) synthesized in Example 9 and 4-methoxyaniline-2-sulfonic acid



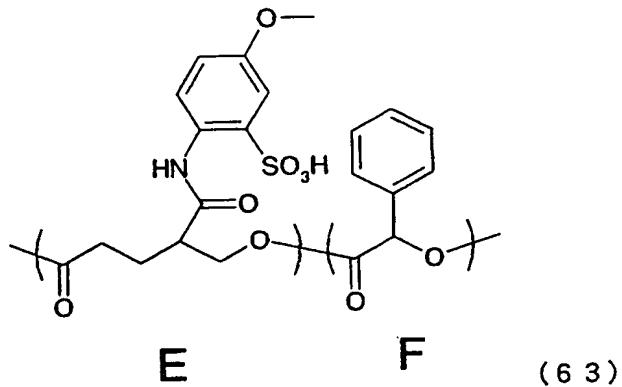
In a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (43) (C: 7 mol%, D: 93 mol%) obtained in Example 9, 0.21 g of 4-methoxyaniline-2-sulfonic acid were added to a 100 ml three-necked flask and agitated with 15.0 ml of pyridine added, and then 0.54 ml of triphenyl phosphite was added and heated at 120°C for 6 hours.

After the reaction ended, re-precipitation was performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N hydrochloric acid for one day and further washed by agitating in water for one day, and 0.32 g of polymer was obtained by carrying out drying under reduced pressure.

The structure of the obtained polymer was determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier transformation-infrared absorption (FT-IR) spectrum

(Nicolet AVATAR360 FT-IR). As a result of performing IR measurement, the peak of $1,695\text{ cm}^{-1}$ resulted from carboxylic acid decreased, and a peak resulted from amide group at $1,658\text{ cm}^{-1}$ was newly observed.

5 The results of $^1\text{H-NMR}$ showed a shift of the peak resulted from the aromatic ring of 4-methoxyaniline-2-sulfonic acid structure and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by
10 10 the following chemical formula (63) as a monomer unit.



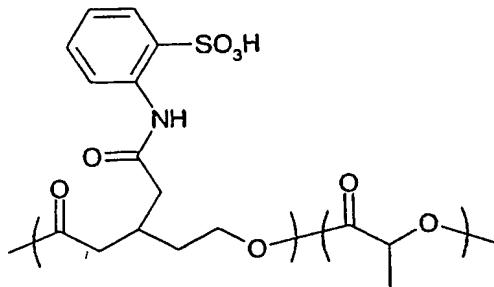
As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (63), it was confirmed that it was a
15 copolymer having a monomer ratio of 7 mol% for unit E and 93 mol% for unit F.

The average molecular weight of the obtained polymer was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr
20

0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, Mn was 6,900, and weight average molecular weight, Mw was 10,100.

Example 30

5 Esterification reaction of the polyhydroxyalkanoate which consists of a unit represented by chemical formula (50) synthesized in Example 16



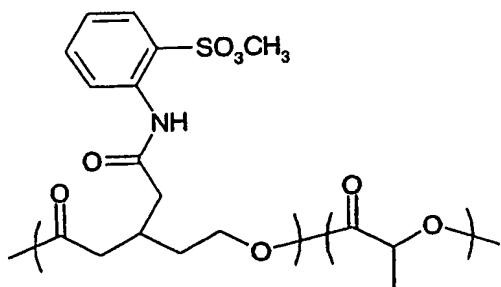
E F (50)

10 0.30 g of the polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (50) (E: 8 mol%, F: 92 mol%) obtained in Example 16 was added to an eggplant flask and dissolved with 21.0 ml of chloroform and 7.0 ml of 15 methanol added and cooled to 0°C. 1.31 ml of 2 mol/L trimethylsilyldiazomethane-hexane solution (product of Aldrich) was added to this, and agitated for 4 hours. After the reaction ended, the solvent was removed with an evaporator and the polymer was 20 collected.

21.0 ml of chloroform, 7.0 ml of methanol were further added to re-dissolve the polymer and the solvent was evaporated by an evaporator. This operation was repeated 3 times.

5 0.30 g of the polymer was obtained by drying the polymer collected here under reduced pressure.

The determination of the structure of the obtained polymer was performed by $^1\text{H-NMR}$ (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature). The results of $^1\text{H-NMR}$ showed a peak resulted from methyl sulfonate at 3 to 4 ppm and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate 10 containing a unit represented by the following 15 chemical formula (64) as a monomer unit.



G

H

(64)

As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical 20 formula (64), it was confirmed that it was a

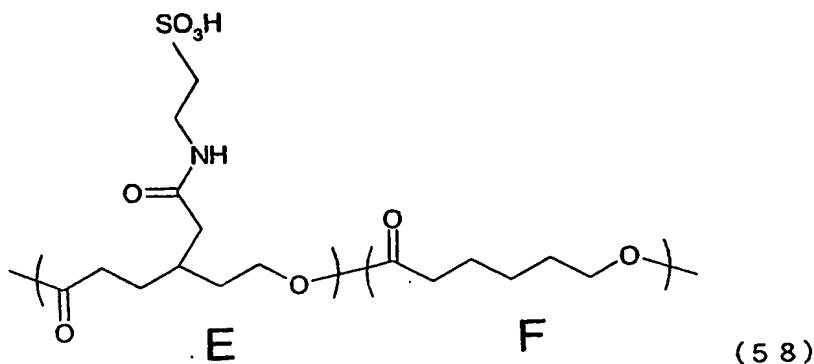
copolymer having a monomer ratio of 8 mol% for unit G and 92 mol% for unit H.

In addition, there was no peak observed resulted from sulfonic acid in acid value titration 5 using Potentiometric Titrator AT510 (product of Kyoto Electronics Manufacturing Co., Ltd.) and it was also made evident from this that sulfonic acid was converted to methyl sulfonate.

The average molecular weight of the obtained 10 polymer was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 15,900, 15 and weight average molecular weight, M_w was 24,200.

Example 31

Esterification reaction of the polyhydroxyalkanoate which consists of a unit represented by chemical formula (58) synthesized in 20 Example 24

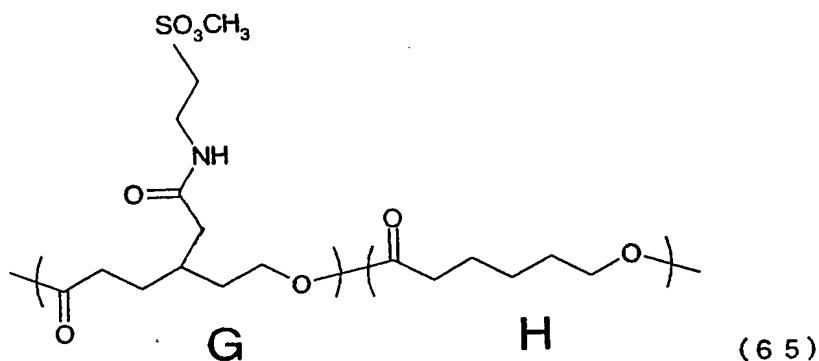


0.30 g of the polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (58) (E: 12 mol%, F: 88 mol%) obtained in 5 Example 24 was added to an eggplant flask and dissolved with 21.0 ml of chloroform and 7.0 ml of methanol added and cooled to 0°C. 1.34 ml of 2 mol/L trimethylsilyldiazomethane-hexane solution (product of Aldrich) was added to this, and agitated for 4 10 hours. After the reaction ended, the solvent was removed with an evaporator and the polymer was collected.

21.0 ml of chloroform, 7.0 ml of methanol were further added to re-dissolve the polymer and the 15 solvent was removed by an evaporator. This operation was repeated 3 times.

0.31 g of the polymer was obtained by drying the polymer collected here under reduced pressure. 20 The determination of the structure of the obtained polymer was performed by ¹H-NMR (FT-NMR:

Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature). The results of $^1\text{H-NMR}$ showed a peak resulted from methyl sulfonate at 3 to 5 4 ppm and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by the following chemical formula (65) as a monomer unit.



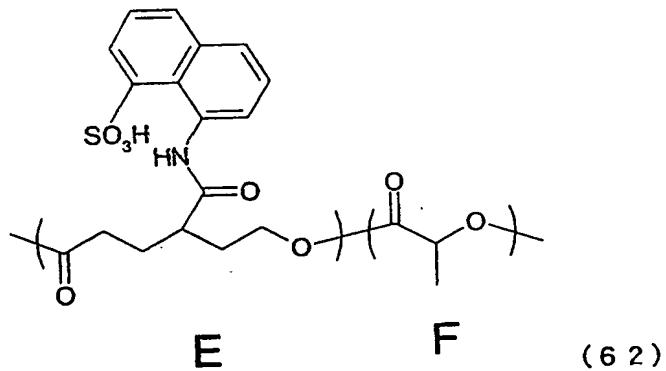
10 As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (65), it was confirmed that it was a copolymer having a monomer ratio of 12 mol% for unit G and 88 mol% for unit H.

15 In addition, there was no peak observed resulted from sulfonic acid in acid value titration using Potentiometric Titrator AT510 (product of Kyoto Electronics Manufacturing Co., Ltd.) and it was also made evident from this that sulfonic acid was 20 converted to methyl sulfonate.

The average molecular weight of the obtained polymer was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 5 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 14,700, and weight average molecular weight, M_w was 21,800.

Example 32

Esterification reaction of the 10 polyhydroxyalkanoate which consists of a unit represented by chemical formula (62) synthesized in Example 28



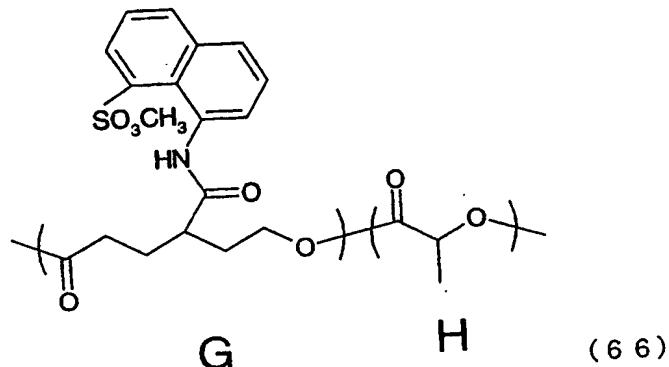
0.30 g of the polyhydroxyalkanoate copolymer 15 which consists of a unit represented by the chemical formula (62) (E: 8 mol%, F: 92 mol%) obtained in Example 28 was added to an eggplant flask and dissolved with 21.0 ml of chloroform and 7.0 ml of methanol added and cooled to 0°C. 1.34 ml of 2 mol/L 20 trimethylsilyldiazomethane-hexane solution (product

of Aldrich) was added to this, and agitated for 4 hours. After the reaction ended, the solvent was removed with an evaporator and the polymer was collected.

5 21.0 ml of chloroform, 7.0 ml of methanol were further added to re-dissolve the polymer and the solvent was removed by an evaporator. This operation was repeated 3 times.

10 0.30 g of the polymer was obtained by drying the polymer collected here under reduced pressure.

The determination of the structure of the obtained polymer was performed by $^1\text{H-NMR}$ (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement 15 temperature: room temperature). The results of $^1\text{H-NMR}$ showed a peak resulted from methyl sulfonate at 3 to 4 ppm and accordingly it was confirmed that the obtained polymer was a polyhydroxyalkanoate containing a unit represented by the following 20 chemical formula (66) as a monomer unit.



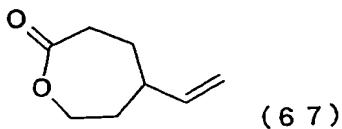
As for the ratio of the units in the polyhydroxyalkanoate represented by the chemical formula (66), it was confirmed that it was a 5 copolymer having a monomer ratio of 8 mol% for unit G and 92 mol% for unit H.

In addition, there was no peak observed resulted from sulfonic acid in acid value titration using Potentiometric Titrator AT510 (product of Kyoto 10 Electronics Manufacturing Co., Ltd.) and it was also made evident from this that sulfonic acid was converted to methyl sulfonate.

The average molecular weight of the obtained polymer was estimated by gel permeation 15 chromatography (GPC; TOSOH HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 7,500, and weight average molecular weight, M_w was 11,400.

Example 33

(Synthesis of polyester using 5-ethenyl-2-oxepanone represented by chemical formula (67) and ϵ -caprolactone)

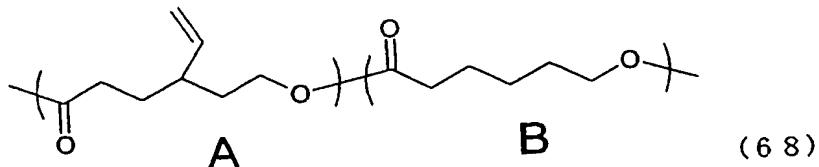


5

0.31 g (2.0 mmol) of 5-ethenyl-2-oxepanone represented by the chemical formula (67), 1.14 g (10.0 mmol) of ϵ -caprolactone, 4.8 ml of 0.01 M toluene solution of tin octylate (tin 2-ethylhexanoate), 4.8 ml of 0.01 M toluene solution of p-tert-butylbenzyl alcohol were placed in a polymerization ampoule and after performing drying under reduced pressure for 1 hour and nitrogen purge, it was heat-sealed under reduced pressure, and heated at 150°C to perform ring-opening polymerization. The reaction was ended after 12 hours and the reaction product was cooled. The obtained polymer was dissolved in chloroform and re-precipitated in methanol in an amount of 10 times of the chloroform required for dissolving the polymer. The precipitation was collected and 0.86 g of polymer was obtained by carrying out drying under reduced pressure.

In order to identify the structure of the

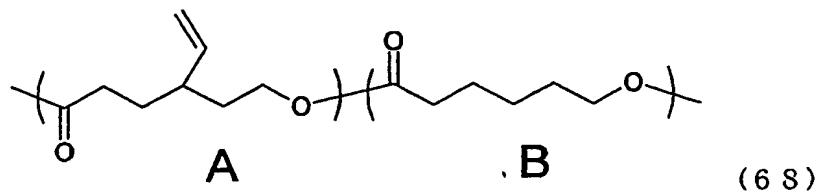
obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate copolymer which contains a unit represented by the following chemical formula (68) as a monomer unit. It was also confirmed that the ratios of monomer units were 12 mol% for A unit and 88 mol% for B unit.



10 The average molecular weight of the obtained
polyhydroxyalkanoate was estimated by gel permeation
chromatography (GPC; TOSOH HLC-8220, column; TOSOH
TSK-GEL Super HM-H, solvent; chloroform, polystyrene
conversion). Consequently, the number average
15 molecular weight, M_n was 17,500, and weight average
molecular weight, M_w was 25,400.

Example 34

Oxidation reaction of polyhydroxyalkanoate which consists of a unit represented by chemical formula (68) synthesized in Example 33

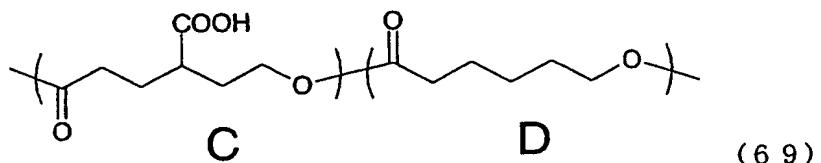


(68)

0.50 g of a polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical formula (68) (A: 12 mol%, B: 88 mol%) obtained in 5 Example 33 was added to an eggplant flask and dissolved with 30 ml of acetone added. This was placed in an ice bath, and 5 ml of acetic acid and 0.40 g of 18-crown-6-ether were added and the mixture was agitated. Next, 0.32 g of potassium permanganate 10 was slowly added on the ice bath, and agitated in an ice bath for 2 hours, and agitated at room temperature for further 18 hours. 60 ml of ethyl acetate was added after the reaction ended, and 45 ml of water was further added. Next, sodium hydrogen 15 sulfite was added until peracid was removed. Then, the pH of the liquid was adjusted to 1 with 1.0 N hydrochloric acid. The organic layer was extracted and washed 3 times with 1.0 N hydrochloric acid. After the organic layer was collected, crude polymer 20 was collected by evaporating the solvent. Next, polymer was collected, after washed with 50 ml of water, 50 ml of methanol and further with 50 ml of water 3 times. Next, it was dissolved in 3 ml of THF,

re-precipitation was performed using methanol in an amount of 50 times of THF required for dissolving the polymer. The precipitation was collected and 0.42 g of polymer was obtained by carrying out drying under reduced pressure.

In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate which contains a unit represented by the following chemical formula (69) as a monomer unit.



The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 14,300, and weight average molecular weight, M_w was 21,500.

In order to calculate the unit of the obtained polyhydroxyalkanoate furthermore, carboxyl groups at the end of the side chains of the

polyhydroxyalkanoate were subjected to methyl esterification using trimethylsilyldiazomethane and the calculation was performed.

30 mg of the target polyhydroxyalkanoate was 5 added to 100 ml-volume eggplant flask and dissolved with 2.1 ml of chloroform and 0.7 ml of methanol added. 0.5 ml of 2 mol/L trimethylsilyldiazomethane-hexane solution was added to this, and agitated at room temperature for 1 hour. After the reaction 10 ended, the solvent was removed and the polymer was collected. It was washed with 50 ml of methanol and the polymer was collected after that. 29 mg of polyhydroxyalkanoate was obtained by carrying out drying under reduced pressure.

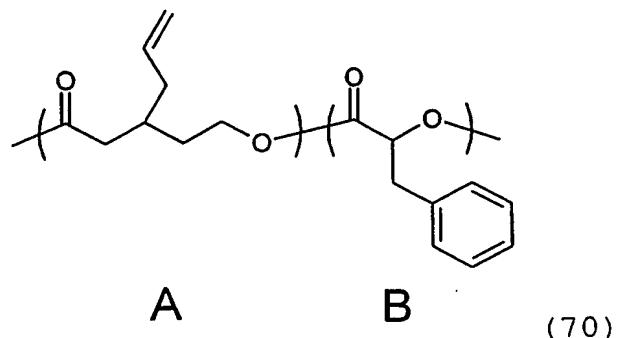
15 NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained polyhydroxyalkanoate represented by the chemical formula (69) was a copolymer having a monomer ratio of 11 mol% for C 20 unit and 89 mol% for D unit.

Example 35

(Synthesis of polyester using tetrahydro-4-(2-propenyl)-2H-pyran-2-one and phenyllactide (3,6-bis(phenylmethyl)-1,4-dioxane-2,5-dione))

25 0.28 g (2.0 mmol) of tetrahydro-4-(2-propenyl)-2H-pyran-2-one represented by the chemical formula (26), 2.96 g (10.0 mmol) of phenyllactide, 4.8 ml of

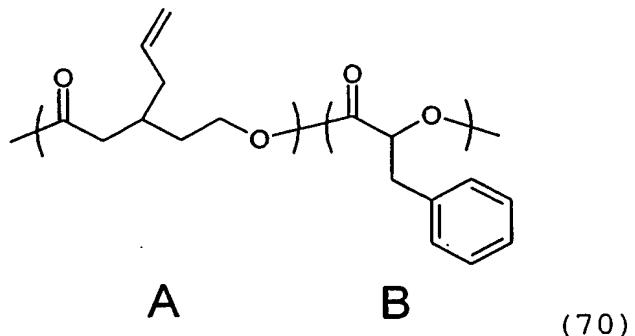
0.01 M toluene solution of tin octylate (tin 2-ethylhexanoate), 4.8 ml of 0.01 M toluene solution of p-tert-butylbenzyl alcohol were placed in a polymerization ampoule and after performing drying 5 under reduced pressure for 1 hour and nitrogen purge, it was heat-sealed under reduced pressure, and heated at 180°C to perform ring-opening polymerization. The reaction was ended after 12 hours and the reaction product was cooled. The obtained polymer was 10 dissolved in chloroform and re-precipitated in methanol in an amount of 10 times of the chloroform required for dissolving the polymer. The precipitation was collected and 2.07 g of polymer was obtained by carrying out drying under reduced 15 pressure. In order to identify the structure of the obtained polymer, NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate copolymer which contains a unit 20 represented by the following chemical formula (70) as a monomer unit. It was also confirmed that the ratios of monomer units were 11 mol% for A unit and 89 mol% for B unit.



The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH 5 TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 20,500, and weight average molecular weight, M_w was 26,600.

Example 36

10 Oxidation reaction of polyhydroxyalkanoate which consists of a unit represented by chemical formula (70) synthesized in Example 35

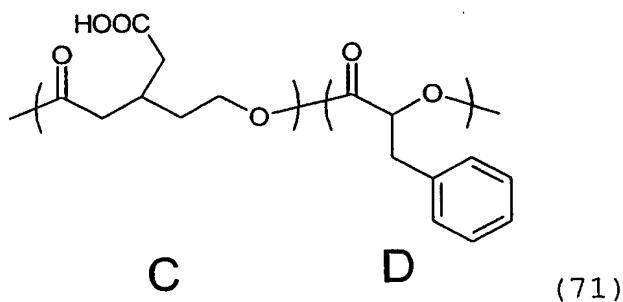


15 0.50 g of a polyhydroxyalkanoate copolymer which consists of a unit represented by the chemical

formula (70) (A: 11 mol%, B: 89 mol%) obtained in Example 35 was added to an eggplant flask and dissolved with 30 ml of acetone added. This was placed in an ice bath, and 5 ml of acetic acid and 5 0.42 g of 18-crown-6-ether were added and the mixture was agitated. Next, 0.30 g of potassium permanganate was slowly added on the ice bath, and agitated in an ice bath for 2 hours, and agitated at room temperature for further 18 hours. 60 ml of ethyl 10 acetate was added after the reaction ended, and 45 ml of water was further added. Next, sodium hydrogen sulfite was added until peracid was removed. Then, the pH of the liquid was adjusted to 1 with 1.0 N hydrochloric acid. The organic layer was extracted 15 and washed 3 times with 1.0 N hydrochloric acid. After the organic layer was collected, crude polymer was collected by evaporating the solvent. Next, polymer was collected, after washed with 50 ml of water, 50 ml of methanol and further with 50 ml of 20 water 3 times. Next, it was dissolved in 3 ml of THF, re-precipitation was performed using methanol in an amount of 50 times of THF required for dissolving the polymer. The precipitation was collected and 0.42 g of polymer was obtained by carrying out drying under 25 reduced pressure.

In order to identify the structure of the obtained polymer, NMR analysis was performed on the

same conditions as in Example 1 and consequently it was confirmed that the obtained compound was a polyhydroxyalkanoate which contains a unit represented by the following chemical formula (71) as 5 a monomer unit.



The average molecular weight of the obtained polyhydroxyalkanoate was estimated by gel permeation chromatography (GPC; TOSOH HLC-8220, column; TOSOH 10 TSK-GEL Super HM-H, solvent; chloroform, polystyrene conversion). Consequently, the number average molecular weight, M_n was 16,800, and weight average molecular weight, M_w was 22,300.

In order to calculate the unit of the obtained 15 polyhydroxyalkanoate furthermore, carboxyl groups at the end of the side chains of the polyhydroxyalkanoate were subjected to methyl esterification using trimethylsilyldiazomethane and the calculation was performed.

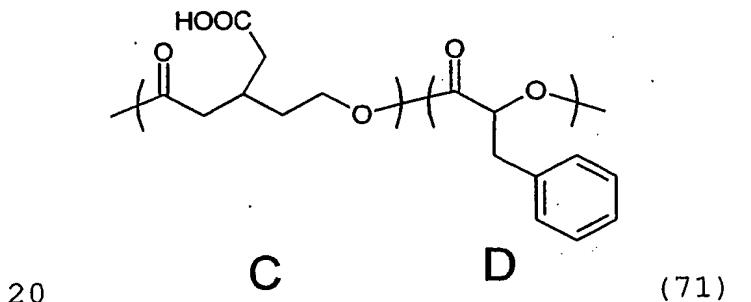
20 30 mg of the target polyhydroxyalkanoate was added to 100 ml-volume eggplant flask and dissolved with 2.1 ml of chloroform and 0.7 ml of methanol

added. 0.5 ml of 2 mol/L trimethylsilyldiazomethane-hexane solution was added to this, and agitated at room temperature for 1 hour. After the reaction ended, the solvent was removed and the polymer was 5 collected. It was washed with 50 ml of methanol and the polymer was collected after that. 29 mg of polyhydroxyalkanoate was obtained by carrying out drying under reduced pressure.

10 NMR analysis was performed on the same conditions as in Example 1 and consequently it was confirmed that the obtained polyhydroxyalkanoate represented by the chemical formula (71) was a copolymer having a monomer ratio of 11 mol% for C unit and 89 mol% for D unit.

15 Example 37

Condensation reaction of polyhydroxyalkanoate which consist of a unit represented by chemical formula (71) synthesized in Example 36 and 2-aminobenzenesulfonic acid



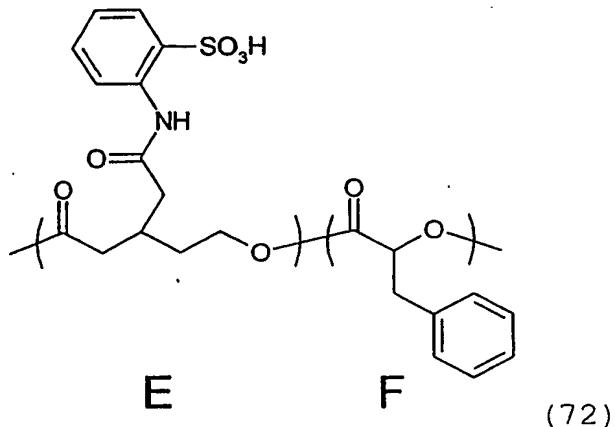
In a nitrogen atmosphere, 0.40 g of the polyhydroxyalkanoate copolymer which consists of a

unit represented by the chemical formula (71) (C: 11 mol%, D: 89 mol%) obtained in Example 36, 0.26 g of 2-aminobenzenesulfonic acid were added to a 100 ml three-necked flask and agitated with 15.0 ml of 5 pyridine added, and then 0.77 ml of triphenyl phosphite was added and heated at 120°C for 6 hours. After the reaction ended, re-precipitation was performed from 150 ml of ethanol to collect the product. The obtained polymer was washed with 1N 10 hydrochloric acid for one day and further washed by agitating in water for one day, and 0.34 g of polymer was obtained by carrying out drying under reduced pressure.

The structure of the obtained polymer was 15 determined by analyzing with $^1\text{H-NMR}$ (FT-NMR: Bruker DPX400; resonant frequency: 400 MHz; nuclide measured: ^1H ; solvent used: d-DMSO; measurement temperature: room temperature) and Fourier transformation-infrared absorption (FT-IR) spectrum 20 (Nicolet AVATAR360 FT-IR). As a result of performing IR measurement, the peak of $1,695\text{ cm}^{-1}$ resulted from carboxylic acid decreased, and a peak resulted from amide group at $1,658\text{ cm}^{-1}$ was newly observed.

The results of $^1\text{H-NMR}$ showed a shift of the peak 25 resulted from the aromatic ring of 2-aminobenzenesulfonic acid structure and accordingly it was confirmed that the obtained polymer was a

polyhydroxyalkanoate containing a unit represented by the following chemical formula (72) as a monomer unit.



As for the ratio of the units in the 5 polyhydroxyalkanoate represented by the chemical formula (72), it was confirmed that it was a copolymer having a monomer ratio of 11 mol% for unit E and 89 mol% for unit F.

The average molecular weight of the obtained 10 polymer was estimated by gel permeation chromatography (GPC; TOSOH HLC-8120, column; Polymer Laboratories PLgel 5 μ MIXED-C, solvent; DMF/LiBr 0.1% (w/v), polystyrene conversion). Consequently, the number average molecular weight, M_n was 16,000, 15 and weight average molecular weight, M_w was 22,700.

A novel polyhydroxyalkanoate which contains a vinyl group, which is a reactive group, on a side chain in a molecule, a novel polyhydroxyalkanoate which contains a carboxyl group in a molecule, a 20 novel polyhydroxyalkanoate which contains a unit

having an amide group and a sulfonic acid group in a molecule, and a production method thereof are provided by the present invention. These new polyhydroxyalkanoates having a vinyl group or a 5 carboxyl group can have functional groups introduced therein utilizing these reactive groups, and accordingly can be applied to functional materials. Furthermore, polyhydroxyalkanoates which contain in a molecule a carboxyl group and a unit having an amide 10 group and a sulfonic acid group are excellent in melt processability, and also excellent in biocompatibility due to their hydrophilicity, and can be expected to be applied to soft materials for medical use, etc.

15

This application claims priority from Japanese Patent Application No. 2004-174784 filed June 11, 2004, which is hereby incorporated by reference 20 herein.